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SUBSTITUTED HETEROCYCLIC COMPOUNDS

Priority is claimed to U.S. Provisional Patent Application Serial No. 60/440,936, filed January 17, 2003, the complete disclosure of which is hereby incorporated by reference.

Field of the Invention

The present invention relates to novel heterocyclic derivatives, in particular piperazine and piperidine derivatives, and to their use in the treatment of various disease states, in particular cardiovascular diseases such as atrial and ventricular arrhythmias, intermittent claudication, Prinzmetal's (variant) angina, stable and unstable angina, exercise induced angina, congestive heart disease, ischemia, reperfusion injury, diabetes, myocardial infarction, and for increasing HDL levels in plasma while lowering LDL levels. The invention also relates to methods for their preparation, and to pharmaceutical compositions containing such compounds.

Background

Certain classes of piperazine compounds are known to be useful for the treatment of cardiovascular diseases, including arrhythmias, angina, myocardial infarction, and related diseases such as intermittent claudication. For example, U.S. Patent No. 4,567,264 discloses a class of substituted piperazine compounds that includes a compound known as ranolazine, (\pm)-N- (2,6-dimethylphenyl)-4-[2-hydroxy-3- (2-methoxyphenoxy)-propyl]-1-piperazineacetamide (RanexaTM), and its pharmaceutically acceptable salts, and their use in the above disease states.

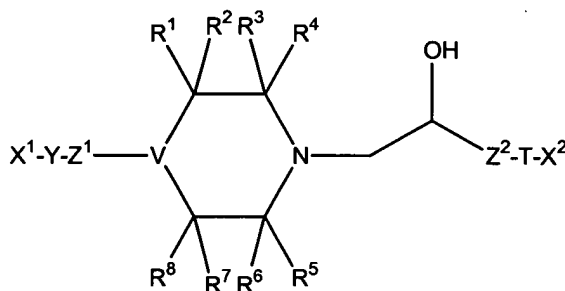
Despite the desirable properties demonstrated by ranolazine, which is a very effective cardiac therapeutic agent, believed to function as a fatty acid oxidation inhibitor,

there remains a need for compounds that have similar therapeutic properties to ranolazine, but are more potent and have a longer half-life.

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SUMMARY OF THE INVENTION

It is an object of this invention to provide novel substituted piperazine and piperidine compounds that are fatty acid oxidation inhibitors with good therapeutic half-lives. Accordingly, in a first aspect, the invention relates to compounds of Formula I:



Formula I

wherein:

- 10 R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are hydrogen, lower alkyl, or -C(O)R;
in which R is -OR⁹ or -NR⁹R¹⁰, where R⁹ and R¹⁰ are hydrogen or lower alkyl; or
R¹ and R², R³ and R⁴, R⁵ and R⁶, R⁷ and R⁸, when taken together with the carbon to
which they are attached, represent carbonyl; or
R¹ and R⁵, or R¹ and R⁷, or R³ and R⁵, or R³ and R⁷, when taken together form a bridging
15 group -(CR¹²R¹³)_n-, in which n is 1, 2 or 3, and R¹² and R¹³ are independently
hydrogen or lower alkyl;
with the proviso that the maximum number of carbonyl groups is 1;
the maximum number of -C(O)NR⁹R¹⁰ groups is 1; and
the maximum number of bridging groups is 1;
20 T is oxygen, sulfur, or NR¹¹, in which R¹¹ is hydrogen or lower alkyl;
V is -N<, -CH<, or -NH-CH<;
X¹ is hydrogen, optionally substituted lower alkyl, optionally substituted cycloalkyl,
optionally substituted aryl, or optionally substituted heteroaryl;
X² is optionally substituted aryl or optionally substituted heteroaryl;

Y is optionally substituted dihydroheteroaryl; and

Z¹ and Z² are independently optionally substituted alkylene of 1-4 carbon atoms.

A second aspect of this invention relates to pharmaceutical formulations, comprising a therapeutically effective amount of a compound of Formula I and at least
5 one pharmaceutically acceptable excipient.

A third aspect of this invention relates to a method of using the compounds of Formula I in the treatment of a disease or condition in a mammal that is amenable to treatment by a fatty acid oxidation inhibitor. Such diseases include, but are not limited to, protection of skeletal muscles against damage resulting from trauma, intermittent
10 claudication, shock, and cardiovascular diseases including atrial and ventricular arrhythmias, Prinzmetal's (variant) angina, unstable angina, exercise induced angina, congestive heart disease, diabetes, myocardial infarction, and for increasing HDL levels in plasma while lowering LDL levels. The compounds of Formula I can also be used to preserve donor tissue and organs used in transplants.

15 A fourth aspect of this invention relates to methods of preparing the compounds of Formula I.

Of the compounds of Formula I, one preferred class includes those compounds in which R¹, R², R³, R⁴, R⁶, R⁷, and R⁸ are hydrogen or methyl and V is -N<. A preferred group within this class includes those compounds in which X¹ is optionally substituted
20 aryl and X² is optionally substituted heteroaryl, particularly where Z¹ is methylene and Z² is methylene or ethylene. One preferred group within the class includes those compounds in which Y is optionally substituted dihydroheteroaryl in which the hetero atoms are chosen from nitrogen and oxygen, particularly where T is oxygen, X¹ is optionally substituted phenyl, Y is optionally substituted dihydroisoxazoliny, and X² is optionally
25 substituted benzothiazolyl. One preferred subgroup includes those compounds in which Y is optionally substituted 5-(4,5-dihydroisoxazol-3-yl). Another preferred subgroup includes those compounds in which Y is optionally substituted 3-(4,5-dihydroisoxazolyl-5-yl).

Of the compounds of Formula I, preferred compounds include
30 (2R)-3-(2-methylbenzothiazol-5-yloxy)-1-{4-[(5-phenyl(4,5-dihydroisoxazol-3-yl))methyl]piperazinyl}propan-2-ol;

(2R)-1-((2S)-2-methyl-4-[(5-phenyl(4,5-dihydroisoxazol-3-yl))methyl]piperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-((3S)-3-methyl-4-[(5-phenyl(4,5-dihydroisoxazol-3-yl))methyl]piperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

5 (2R)-3-(2-methylbenzothiazol-5-yloxy)-1-(4-[[5-(2,4,6-trimethylphenyl)(4,5-dihydroisoxazol-3-yl)]methyl]piperazinyl)propan-2-ol;

(2R)-1-((3S)-4-[[5-(2,6-dimethylphenyl)(4,5-dihydroisoxazol-3-yl)]methyl]-3-methylpiperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-(4-[[[(5R)-3-(4-fluorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl](2S)-2-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

10 (2R)-1-(4-[[[(5R)-3-(4-fluorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl]piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-(4-[[[(5R)-3-(4-fluorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl](3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

15 (2R)-1-[4-(([(5R)-3-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)piperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[4-(([(5R)-3-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)(3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[4-(([(5R)-3-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)(2S)-2-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

20 (2R)-1-[4-(([(5R)-3-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)piperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[4-(([(5R)-3-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)(3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

25 (2R)-1-[4-(([(5R)-3-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)(2S)-2-methylpiperazinyl]-3-benzothiazol-5-yloxypropan-2-ol;

(2R)-1-{4-[[[(4S)-2-phenyl(1,3-oxazolin-4-yl))methyl]piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-((3S)-4-[[[(4S)-2-phenyl(1,3-oxazolin-4-yl))methyl]-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

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(2R)-1-{4-[[[(5R)-3-phenyl(4,5-dihydroisoxazol-5-yl)methyl](2S)-2-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-{4-[[[(4S,5S)-2-methyl-5-phenyl(1,3-oxazolin-4-yl)methyl]piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

5 (2R)-3-{(2S)-4-[[[(4S,5S)-2-methyl-5-phenyl(1,3-oxazolin-4-yl)methyl]-2-methylpiperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[(2S)-2-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

10 (2R)-3-(2-methylbenzothiazol-5-yloxy)-1-[4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-[(3S)-3-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[(2S)-2-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

15 (2R)-1-{4-[[[(5R)-3-phenyl(4,5-dihydroisoxazol-5-yl)methyl](3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-{4-[[[(5R)-3-phenyl(4,5-dihydroisoxazol-5-yl)methyl]piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

20 (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[2-chlorophenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

25 (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[(2S)-2-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[(2S)-2-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

30 (2R)-3-[(3S)-3-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-[(3S)-3-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-(4-{[2-(4-fluorophenyl)(1,3-oxazolin-4-yl)]methyl}piperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

5 (2R)-3-((2S)-2-methyl-4-{[3-(4-methylphenyl)(4,5-dihydroisoxazol-5-yl)]methyl}piperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol; and

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-(4-{[3-(4-chlorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl}(2S)-2-methylpiperazinyl) propan-2-ol.

More preferred compounds include:

10 (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-[(3S)-3-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

15 (2R)-3-[(3S)-3-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-(2-methylbenzothiazol-5-yloxy)-1-[4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-[4-({(5R)-3-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

20 (2R)-1-{4-[[((5R)-3-phenyl(4,5-dihydroisoxazol-5-yl))methyl](2S)-2-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-(2-methylbenzothiazol-5-yloxy)-1-{4-[(5-phenyl(4,5-dihydroisoxazol-3-yl))methyl]piperazinyl}propan-2-ol;

25 (2R)-1-(4-{[(5R)-3-(4-fluorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl}(3S)-3-methylpiperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[(2S)-2-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol; and

(2R)-1-[4-({(5R)-3-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol.

Definitions and General Parameters

The term “alkyl” refers to a monoradical branched or unbranched saturated hydrocarbon chain having from 1 to 20 carbon atoms. This term is exemplified by groups such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, t-butyl, n-hexyl, n-decyl, tetradecyl, and the like.

The term “substituted alkyl” refers to:

- 1) an alkyl group as defined above, having from 1 to 5 substituents, preferably 1 to 3 substituents, selected from the group consisting of alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxy-carbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy, heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocycloxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2; or
- 2) an alkyl group as defined above that is interrupted by 1-5 atoms or groups independently chosen from oxygen, sulfur and -NR_a-, where R_a is chosen from hydrogen, alkyl, cycloalkyl, alkenyl, cycloalkenyl, alkynyl, aryl, heteroaryl and heterocyclyl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2; or
- 3) an alkyl group as defined above that has both from 1 to 5 substituents as defined above and is also interrupted by 1-5 atoms or groups as defined above.

The term “lower alkyl” refers to a monoradical branched or unbranched saturated hydrocarbon chain having from 1 to 6 carbon atoms. This term is exemplified by groups

such as methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, t-butyl, n-hexyl, and the like.

The term “substituted lower alkyl” refers to lower alkyl as defined above having 1 to 5 substituents, preferably 1 to 3 substituents, as defined for substituted alkyl, or a lower alkyl group as defined above that is interrupted by 1-5 atoms as defined for substituted alkyl, or a lower alkyl group as defined above that has both from 1 to 5 substituents as defined above and is also interrupted by 1-5 atoms as defined above.

The term “alkylene” refers to a diradical of a branched or unbranched saturated hydrocarbon chain, preferably having from 1 to 20 carbon atoms, preferably 1-10 carbon atoms, more preferably 1-6 carbon atoms. This term is exemplified by groups such as methylene (-CH₂-), ethylene (-CH₂CH₂-), the propylene isomers (e.g., -CH₂CH₂CH₂- and -CH(CH₃)CH₂-) and the like.

The term “lower alkylene” refers to a diradical of a branched or unbranched saturated hydrocarbon chain, preferably having from 1 to 6 carbon atoms.

The term “substituted alkylene” refers to:

- (1) an alkylene group as defined above having from 1 to 5 substituents selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxycarbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy, heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocycloxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2; or
- (2) an alkylene group as defined above that is interrupted by 1-5 atoms or groups independently chosen from oxygen, sulfur and NR_a-, where R_a is chosen from hydrogen, optionally substituted alkyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl

and heterocyclyl, or groups selected from carbonyl, carboxyester, carboxyamide and sulfonyl; or

- (3) an alkylene group as defined above that has both from 1 to 5 substituents as defined above and is also interrupted by 1-20 atoms as defined above. Examples of substituted alkylenes are chloromethylene (-CH(Cl)-), aminoethylene (-CH(NH₂)CH₂-), methylaminoethylene (-CH(NHMe)CH₂-), 2-carboxypropylene isomers(-CH₂CH(CO₂H)CH₂-), ethoxyethyl (-CH₂CH₂O-CH₂CH₂-), ethylmethylaminoethyl (-CH₂CH₂N(CH₃)CH₂CH₂-), 1-ethoxy-2-(2-ethoxy-ethoxy)ethane (-CH₂CH₂O-CH₂CH₂-OCH₂CH₂-OCH₂CH₂-), and the like.

The term “aralkyl” refers to an aryl group covalently linked to an alkylene group, where aryl and alkylene are defined herein. “Optionally substituted aralkyl” refers to an optionally substituted aryl group covalently linked to an optionally substituted alkylene group. Such aralkyl groups are exemplified by benzyl, phenylethyl, 3-(4-methoxyphenyl)propyl, and the like.

The term “alkoxy” refers to the group R-O-, where R is optionally substituted alkyl or optionally substituted cycloalkyl, or R is a group -Y-Z, in which Y is optionally substituted alkylene and Z is optionally substituted alkenyl, optionally substituted alkynyl; or optionally substituted cycloalkenyl, where alkyl, alkenyl, alkynyl, cycloalkyl and cycloalkenyl are as defined herein. Preferred alkoxy groups are optionally substituted alkyl-O- and include, by way of example, methoxy, ethoxy, n-propoxy, isopropoxy, n-butoxy, tert-butoxy, sec-butoxy, n-pentoxy, n-hexoxy, 1,2-dimethylbutoxy, trifluoromethoxy, and the like.

The term “alkylthio” refers to the group R-S-, where R is as defined for alkoxy.

The term “alkenyl” refers to a monoradical of a branched or unbranched unsaturated hydrocarbon group preferably having from 2 to 20 carbon atoms, more preferably 2 to 10 carbon atoms and even more preferably 2 to 6 carbon atoms and having 1-6, preferably 1, double bond (vinyl). Preferred alkenyl groups include ethenyl or vinyl (-CH=CH₂), 1-propylene or allyl (-CH₂CH=CH₂), isopropylene (-C(CH₃)=CH₂), bicyclo[2.2.1]heptene, and the like. In the event that alkenyl is attached to nitrogen, the double bond cannot be alpha to the nitrogen.

The term “lower alkenyl” refers to alkenyl as defined above having from 2 to 6

carbon atoms.

The term “substituted alkenyl” refers to an alkenyl group as defined above having from 1 to 5 substituents, and preferably 1 to 3 substituents, selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxycarbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy, heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocyclooxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

The term “alkynyl” refers to a monoradical of an unsaturated hydrocarbon, preferably having from 2 to 20 carbon atoms, more preferably 2 to 10 carbon atoms and even more preferably 2 to 6 carbon atoms and having at least 1 and preferably from 1-6 sites of acetylene (triple bond) unsaturation. Preferred alkynyl groups include ethynyl, (-C≡CH), propargyl (or prop-1-yn-3-yl, -CH₂C≡CH), and the like. In the event that alkynyl is attached to nitrogen, the triple bond cannot be alpha to the nitrogen.

The term “substituted alkynyl” refers to an alkynyl group as defined above having from 1 to 5 substituents, and preferably 1 to 3 substituents, selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxycarbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy, heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocyclooxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

The term “aminocarbonyl” refers to the group -C(O)NRR where each R is independently hydrogen, alkyl, aryl, heteroaryl, heterocyclyl or where both R groups are joined to form a heterocyclic group (e.g., morpholino). Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents
5 chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

The term “acylamino” refers to the group -NRC(O)R where each R is independently hydrogen, alkyl, aryl, heteroaryl, or heterocyclyl. Unless otherwise
10 constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

The term “acyloxy” refers to the groups -O(O)C-alkyl, -O(O)C-cycloalkyl, -
15 O(O)C-aryl, -O(O)C-heteroaryl, and -O(O)C-heterocyclyl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

20 The term “aryl” refers to an aromatic carbocyclic group of 6 to 20 carbon atoms having a single ring (e.g., phenyl) or multiple rings (e.g., biphenyl), or multiple condensed (fused) rings (e.g., naphthyl or anthryl). Preferred aryls include phenyl, naphthyl and the like.

Unless otherwise constrained by the definition for the aryl substituent, such aryl
25 groups can optionally be substituted with from 1 to 5 substituents, preferably 1 to 3 substituents, selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxycarbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy,
30 heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocyclooxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl,

-SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl
5 and n is 0, 1 or 2.

The term “aryloxy” refers to the group aryl-O- wherein the aryl group is as defined above, and includes optionally substituted aryl groups as also defined above. The term “arylthio” refers to the group R-S-, where R is as defined for aryl.

The term “amino” refers to the group -NH₂.

10 The term “substituted amino” refers to the group -NRR where each R is independently selected from the group consisting of hydrogen, alkyl, cycloalkyl, carboxyalkyl (for example, benzyloxycarbonyl), aryl, heteroaryl and heterocyclyl provided that both R groups are not hydrogen, or a group -Y-Z, in which Y is optionally substituted alkylene and Z is alkenyl, cycloalkenyl, or alkynyl. Unless otherwise
15 constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

The term “carboxyalkyl” refers to the groups -C(O)O-alkyl,
20 -C(O)O-cycloalkyl, where alkyl and cycloalkyl, are as defined herein, and may be optionally further substituted by alkyl, alkenyl, alkynyl, alkoxy, halogen, CF₃, amino, substituted amino, cyano, or -S(O)_nR, in which R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

The term “cycloalkyl” refers to cyclic alkyl groups of from 3 to 20 carbon atoms
25 having a single cyclic ring or multiple condensed rings. Such cycloalkyl groups include, by way of example, single ring structures such as cyclopropyl, cyclobutyl, cyclopentyl, cyclooctyl, and the like, or multiple ring structures such as adamantanyl, and bicyclo[2.2.1]heptane, or cyclic alkyl groups to which is fused an aryl group, for example indan, and the like.

30 The term “substituted cycloalkyl” refers to cycloalkyl groups having from 1 to 5 substituents, and preferably 1 to 3 substituents, selected from the group consisting of

alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxycarbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy, heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocyclooxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2.

The term “halogen” or “halo” refers to fluoro, bromo, chloro, and iodo.

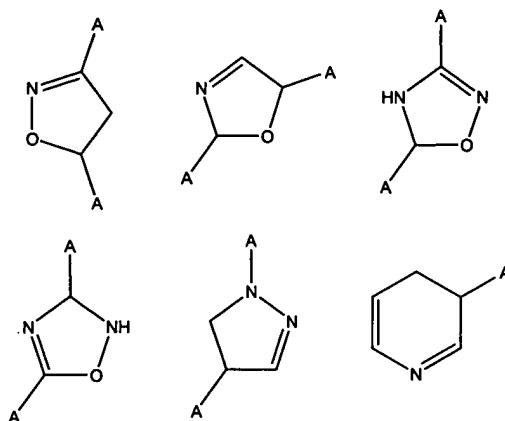
The term “acyl” denotes a group -C(O)R, in which R is hydrogen, optionally substituted alkyl, optionally substituted cycloalkyl, optionally substituted heterocyclyl, optionally substituted aryl, and optionally substituted heteroaryl.

The term “heteroaryl” refers to an aromatic group (i.e., unsaturated) comprising 1 to 15 carbon atoms and 1 to 4 heteroatoms selected from oxygen, nitrogen and sulfur within at least one ring.

Unless otherwise constrained by the definition for the heteroaryl substituent, such heteroaryl groups can be optionally substituted with 1 to 5 substituents, preferably 1 to 3 substituents selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxycarbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy, heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocyclooxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2. Such heteroaryl groups can have a single ring (e.g., pyridyl, furyl, oxadiazolyl, oxazolyl, isoxazolyl, pyrazolyl) or multiple condensed rings (e.g., bicyclic

heteroaryl groups, such as indoliziny, benzothiazoly, benzoxazoly, benzothienyl, and the like). Examples of nitrogen heterocycles and heteroaryls include, but are not limited to, pyrrole, imidazole, pyrazole, pyridine, pyrazine, pyrimidine, pyridazine, indolizine, isoindole, indole, indazole, purine, quinolizine, isoquinoline, quinoline, phthalazine, naphthylpyridine, quinoxaline, quinazoline, cinnoline, pteridine, carbazole, carboline, phenanthridine, acridine, phenanthroline, thiazole, isothiazole, phenazine, isoxazole, phenoxazine, phenothiazine, imidazolidine, imidazoline, and the like as well as N-alkoxy-nitrogen containing heteroaryl compounds.

The term “dihydroheteroaryl” refers to a heteroaryl group as defined above in which one double bond has been saturated, and one double bond remains unsaturated. That is, a partially saturated heteroaryl group. Examples of such groups are:



in which A represents the points of attachment, which are derivatives of; 3,5-(4,5-dihydroisoxazole), 2,5-(1,3-oxazoline), 3,5-(2,3-dihydro-1,2,4-oxadiazole), 3,5-(4,5-dihydro-1,2,4-oxadiazole), 1,4-pyrazoline, 4,5-dihydropyridine, and the like.

Unless otherwise constrained by the definition for the dihydroheteroaryl substituent, such dihydroheteroaryl groups can be optionally substituted in the same manner as heteroaryl.

The term “heteroaryloxy” refers to the group heteroaryl-O-.

The term “heterocyclyl” refers to a monoradical saturated or partially unsaturated group having a single ring or multiple condensed rings, having from 1 to 40 carbon atoms and from 1 to 10 hetero atoms, preferably 1 to 4 heteroatoms, selected from nitrogen, sulfur, phosphorus, and/or oxygen within the ring.

Unless otherwise constrained by the definition for the heterocyclic substituent, such heterocyclic groups can be optionally substituted with 1 to 5, and preferably 1 to 3 substituents, selected from the group consisting of alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl, cycloalkenyl, acyl, acylamino, acyloxy, amino, aminocarbonyl, alkoxycarbonylamino, azido, cyano, halogen, hydroxy, keto, thiocarbonyl, carboxy, carboxyalkyl, arylthio, heteroarylthio, heterocyclylthio, thiol, alkylthio, aryl, aryloxy, heteroaryl, aminosulfonyl, aminocarbonylamino, heteroaryloxy, heterocyclyl, heterocycloxy, hydroxyamino, alkoxyamino, nitro, -SO-alkyl, -SO-aryl, -SO-heteroaryl, -SO₂-alkyl, SO₂-aryl and -SO₂-heteroaryl. Unless otherwise constrained by the definition, all substituents may optionally be further substituted by 1-3 substituents chosen from alkyl, carboxy, carboxyalkyl, aminocarbonyl, hydroxy, alkoxy, halogen, CF₃, amino, substituted amino, cyano, and -S(O)_nR, where R is alkyl, aryl, or heteroaryl and n is 0, 1 or 2. Heterocyclic groups can have a single ring or multiple condensed rings. Preferred heterocyclics include tetrahydrofuranyl, morpholino, piperidinyl, and the like.

The term “thiol” refers to the group -SH.

The term “substituted alkylthio” refers to the group -S-substituted alkyl.

The term “heteroarylthiol” refers to the group -S-heteroaryl wherein the heteroaryl group is as defined above including optionally substituted heteroaryl groups as also defined above.

The term “sulfoxide” refers to a group -S(O)R, in which R is alkyl, aryl, or heteroaryl. “Substituted sulfoxide” refers to a group -S(O)R, in which R is substituted alkyl, substituted aryl, or substituted heteroaryl, as defined herein.

The term “sulfone” refers to a group -S(O)₂R, in which R is alkyl, aryl, or heteroaryl. “Substituted sulfone” refers to a group -S(O)₂R, in which R is substituted alkyl, substituted aryl, or substituted heteroaryl, as defined herein.

The term “keto” refers to a group -C(O)-. The term “thiocarbonyl” refers to a group -C(S)-. The term “carboxy” refers to a group -C(O)-OH.

“Optional” or “optionally” means that the subsequently described event or circumstance may or may not occur, and that the description includes instances where said event or circumstance occurs and instances in which it does not.

The term "compound of Formula I" is intended to encompass the compounds of the invention as disclosed, and the pharmaceutically acceptable salts, pharmaceutically acceptable esters, and polymorphs and prodrugs of such compounds. Additionally, the compounds of the invention may possess one or more asymmetric centers, and can be produced as a racemic mixture or as individual enantiomers or diastereoisomers. The number of stereoisomers present in any given compound of Formula I depends upon the number of asymmetric centers present (there are 2^n stereoisomers possible where n is the number of asymmetric centers). The individual stereoisomers may be obtained by resolving a racemic or non-racemic mixture of an intermediate at some appropriate stage of the synthesis, or by resolution of the compound of Formula I by conventional means. The individual stereoisomers (including individual enantiomers and diastereoisomers) as well as racemic and non-racemic mixtures of stereoisomers are encompassed within the scope of the present invention, all of which are intended to be depicted by the structures of this specification unless otherwise specifically indicated.

"Isomers" are different compounds that have the same molecular formula.

"Stereoisomers" are isomers that differ only in the way the atoms are arranged in space.

"Enantiomers" are a pair of stereoisomers that are non-superimposable mirror images of each other. A 1:1 mixture of a pair of enantiomers is a "racemic" mixture. The term "(±)" is used to designate a racemic mixture where appropriate.

"Diastereoisomers" are stereoisomers that have at least two asymmetric atoms, but which are not mirror-images of each other.

The absolute stereochemistry is specified according to the Cahn-Ingold-Prelog R-S system. When the compound is a pure enantiomer the stereochemistry at each chiral carbon may be specified by either R or S. Resolved compounds whose absolute configuration is unknown are designated (+) or (-) depending on the direction (dextro- or laevorotary) which they rotate the plane of polarized light at the wavelength of the sodium D line.

The term "therapeutically effective amount" refers to that amount of a compound of Formula I that is sufficient to effect treatment, as defined below, when administered to a mammal in need of such treatment. The therapeutically effective amount will vary depending upon the subject and disease condition being treated, the weight and age of the

subject, the severity of the disease condition, the manner of administration and the like, which can readily be determined by one of ordinary skill in the art.

The term “treatment” or “treating” means any treatment of a disease in a mammal, including:

- 5 (i) preventing the disease, that is, causing the clinical symptoms of the disease not to develop;
- (ii) inhibiting the disease, that is, arresting the development of clinical symptoms; and/or
- (iii) relieving the disease, that is, causing the regression of clinical symptoms.

10 In many cases, the compounds of this invention are capable of forming acid and/or base salts by virtue of the presence of amino and/or carboxyl groups or groups similar thereto. The term “pharmaceutically acceptable salt” refers to salts that retain the biological effectiveness and properties of the compounds of Formula I, and which are not biologically or otherwise undesirable. Pharmaceutically acceptable base addition salts can
15 be prepared from inorganic and organic bases. Salts derived from inorganic bases, include by way of example only, sodium, potassium, lithium, ammonium, calcium and magnesium salts. Salts derived from organic bases include, but are not limited to, salts of primary, secondary and tertiary amines, such as alkyl amines, dialkyl amines, trialkyl amines, substituted alkyl amines, di(substituted alkyl) amines, tri(substituted alkyl)
20 amines, alkenyl amines, dialkenyl amines, trialkenyl amines, substituted alkenyl amines, di(substituted alkenyl) amines, tri(substituted alkenyl) amines, cycloalkyl amines, di(cycloalkyl) amines, tri(cycloalkyl) amines, substituted cycloalkyl amines, disubstituted cycloalkyl amine, trisubstituted cycloalkyl amines, cycloalkenyl amines, di(cycloalkenyl) amines, tri(cycloalkenyl) amines, substituted cycloalkenyl amines, disubstituted
25 cycloalkenyl amine, trisubstituted cycloalkenyl amines, aryl amines, diaryl amines, triaryl amines, heteroaryl amines, diheteroaryl amines, triheteroaryl amines, heterocyclic amines, diheterocyclic amines, triheterocyclic amines, mixed di- and tri-amines where at least two of the substituents on the amine are different and are selected from the group consisting of alkyl, substituted alkyl, alkenyl, substituted alkenyl, cycloalkyl, substituted
30 cycloalkyl, cycloalkenyl, substituted cycloalkenyl, aryl, heteroaryl, heterocyclic, and the like. Also included are amines where the two or three substituents, together with the

amino nitrogen, form a heterocyclic or heteroaryl group.

Specific examples of suitable amines include, by way of example only, isopropylamine, trimethyl amine, diethyl amine, tri(iso-propyl) amine, tri(n-propyl) amine, ethanolamine, 2-dimethylaminoethanol, tromethamine, lysine, arginine, histidine, 5 caffeine, procaine, hydrabamine, choline, betaine, ethylenediamine, glucosamine, N-alkylglucamines, theobromine, purines, piperazine, piperidine, morpholine, N-ethylpiperidine, and the like.

Pharmaceutically acceptable acid addition salts may be prepared from inorganic and organic acids. Salts derived from inorganic acids include hydrochloric acid, 10 hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like. Salts derived from organic acids include acetic acid, propionic acid, glycolic acid, pyruvic acid, oxalic acid, malic acid, malonic acid, succinic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, p-toluene-sulfonic acid, salicylic acid, and the like.

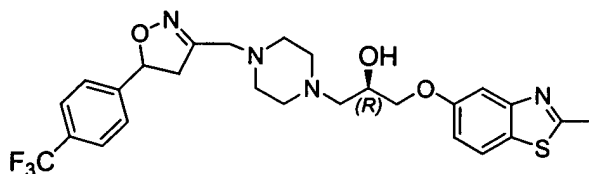
15 As used herein, “pharmaceutically acceptable carrier” includes any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents and the like. The use of such media and agents for pharmaceutically active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active ingredient, its use in the 20 therapeutic compositions is contemplated. Supplementary active ingredients can also be incorporated into the compositions.

“Fatty acid oxidation inhibitors” refers to compounds that suppress ATP production from the oxidation of fatty acids and consequently stimulate ATP production from the oxidation of glucose and lactate. In the heart, most of the ATP production is 25 acquired through the metabolism of fatty acids. The metabolism of glucose and lactate provides a lesser proportion of ATP. However, the generation of ATP from fatty acids is less efficient with respect to oxygen consumption than the generation of ATP from the oxidation of glucose and lactate. Thus, the use of fatty acid oxidation inhibitors results in more energy production per molecule of oxygen consumed, allowing the heart to be 30 energized more efficiently. Fatty acid oxidation inhibitors are especially useful, therefore, for treating an ischemic environment in which oxygen levels are reduced.

Nomenclature

The naming and numbering of the compounds of the invention is illustrated with a representative compound of Formula I in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are hydrogen, T is oxygen, X¹ is (4-trifluoromethyl)phenyl, X² is 2-methylbenzothiazol-5-yl,

5 Y is 5-(4,5-dihydroisoxazol-3-yl), and Z¹ and Z² are methylene:



which is named:

(2R)-3-(2-methylbenzothiazol-5-yloxy)-1-[4-({5-[4-(trifluoromethyl)phenyl](4,5-
10 dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

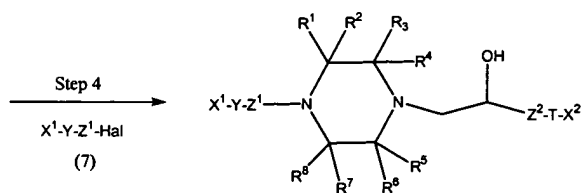
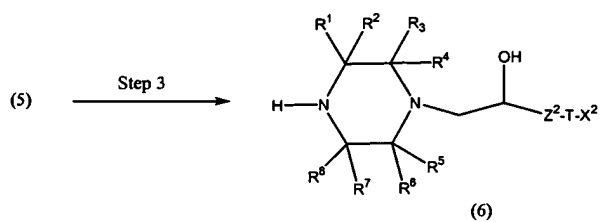
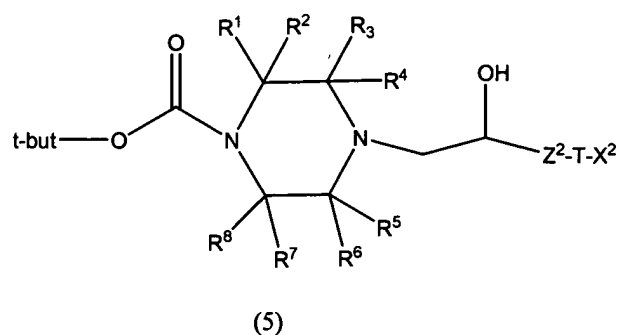
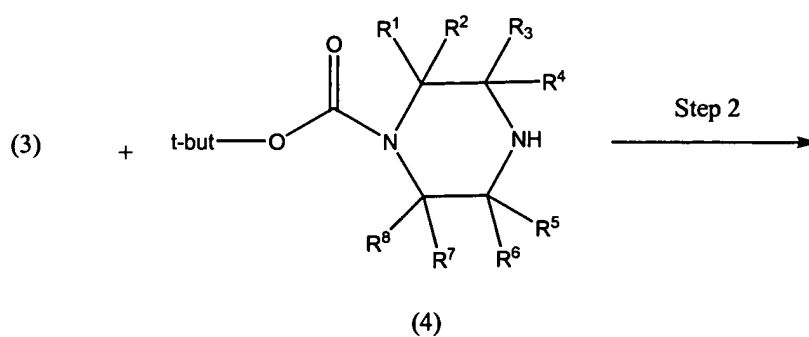
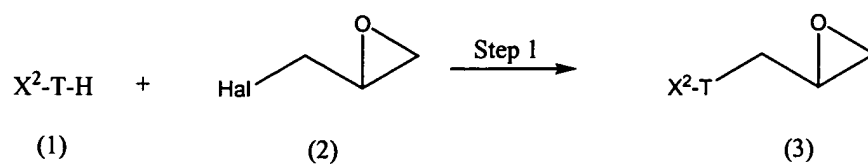
or alternatively;

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[4-(trifluoromethyl)phenyl](4,5-
dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol.

Synthesis of the Compounds of Formula I

One method of preparing the compounds of Formula I is shown in Reaction
Scheme I.

REACTION SCHEME I



in which $\text{R}^1, \text{R}^2, \text{R}^3, \text{R}^4, \text{R}^5, \text{R}^6, \text{R}^7, \text{R}^8, \text{T}, \text{X}^1, \text{X}^2, \text{Y},$ and Z^1 , are as defined in the Summary of the Invention, Z^2 is methylene, Hal is halogen, and t-but is tertiary butyl.

Starting Materials

The compounds of formula (1), (2), and (3) are either commercially available or can be made by conventional methods well known to those of ordinary skill in the art.

For example, the precursor to a compound of formula (4) where R¹ and R⁵ when taken together represent a bridging methylene group, i.e.;



is commercially available [(1S,4S)-(+)-2,5-diazabicyclo[2.2.1]heptane], or can be made by a procedure disclosed in J. Org. Chem., 1990, 55, 1684-7. Similarly, the precursor to a compound of formula (4) where R¹ and R⁵ when taken together represent a bridging ethylene group, and the precursor to a compound of formula (4) where R¹ and R⁷ when taken together represent a bridging ethylene group, can be made by published procedures found in J. Med. Chem., 1974, 17, 481-7. The precursor to a compound of formula (4) in which R¹, R², R³, R⁴, R⁵, R⁶, and R⁷ are hydrogen and R⁸ is -C(O)NH₂ is prepared from piperazine-2-carboxamide, a commercially available compound.

Step 1 - Preparation of Formula (3)

The compound of formula (3) is prepared conventionally by reaction of a compound of formula (1), for example 5-hydroxy-2-methylbenzothiazole, with an epoxide of formula (2), which may be racemic or chiral. In general, the two compounds are mixed in an inert solvent, preferably a ketone, for example acetone, and a tertiary organic base or an inorganic base, preferably potassium carbonate, at a temperature of about reflux, for about 8-48 hours, preferably overnight. When the reaction is substantially complete, the product of formula (3) is isolated by conventional means, for example by filtration, removal of the solvent under reduced pressure, followed by chromatography of the residue on silica gel. Alternatively, after filtration the product can be crystallized from the filtrate.

Step 2 - Preparation of Formula (5)

The compound of formula (3) is then reacted with a protected piperazine of formula (4). In general, the two compounds are mixed in an inert solvent, preferably a halogenated solvent, for example methylene chloride, optionally in the presence of a catalyst, for example ytterbium (III) trifluoromethanesulfonate. In the presence of a catalyst the reaction is conducted at about 0-30°C, preferably at about room temperature, for about 8-48 hours, preferably overnight. In the absence of a catalyst, the mixture is refluxed for a similar period of time in ethanol in the presence of triethylamine. When the reaction is substantially complete, the product of formula (5) is isolated by conventional means, for example by removal of the solvent under reduced pressure, followed by chromatography of the residue on silica gel.

Step 3 - Preparation of Formula (6)

The compound of formula (5) is then deprotected by hydrolyzing the t-butyl ester. In general, the compound of formula (5) is dissolved in a mixture of an inert solvent, preferably a halogenated solvent, for example methylene chloride, and a strong acid, for example trifluoroacetic acid. The reaction is conducted at about 0-30°C, preferably at about room temperature, for about 8-48 hours, preferably overnight. When the reaction is substantially complete, the product of formula (6) is isolated by conventional means, for example by adding a base to remove excess acid, and removal of the solvent under reduced pressure.

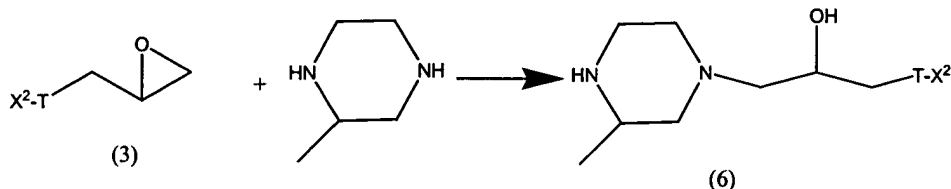
Step 4 - Preparation of a Compound of Formula I

The compound of formula (6) is then reacted with a compound of formula (7) (X^1-Y-Z^1-Hal), for example 3-(chloromethyl)-5-[4-(trifluoromethyl)phenyl]-4,5-dihydroisoxazole. Examples of such compounds are 3-(4-trifluoromethylphenyl)-5-chloromethyl-2,3 dihydro-1,2,4-oxadiazole, 5-bromomethyl-4-methyl-2-phenyl-1,2,3-triazoline, or 4-chloromethyl-2-(4-trifluoromethylphenyl)-1,3-thiazoline, and the like. Such compounds are either commercially available, prepared by means well known in the art (see, for example, see J. Med. Chem, 1996, 39, 237-243) or prepared as shown herein. In general, the two compounds are mixed in an inert solvent, preferably a protic solvent,

for example ethanol, in the presence of an inorganic or tertiary organic base, preferably triethylamine. The reaction is conducted at about 30-100°C, preferably at about reflux, for about 8-48 hours, preferably overnight. When the reaction is substantially complete, the product of Formula I is isolated by conventional means, for example by removal of the solvent under reduced pressure, followed by chromatography.

A modified procedure is preferred for preparing compounds of Formula I in which R⁸ is lower alkyl and R¹-R⁷ are hydrogen. An example where R⁸ is methyl is shown in Reaction Scheme IA.

REACTION SCHEME IA



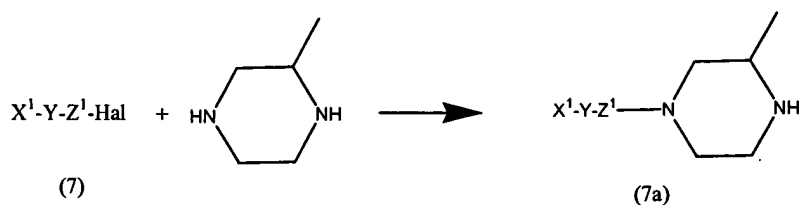
Step 1 - Preparation of Formula (6)

The compound of formula (3) is reacted with 2-methylpiperazine. In general, the two compounds are mixed in a protic solvent, for example ethanol. The reaction is conducted at about 5-100°C, preferably at about 80°C, for about 1-12 hours, preferably about 5 hours. When the reaction is substantially complete, the product of formula (6) is isolated by conventional means, for example by removal of the solvent under reduced pressure, followed by chromatography of the residue on silica gel.

The compound of formula (6) is then reacted with a compound of formula (7) as described above in Reaction Scheme I, step 4, to provide a compound of Formula I in which R¹, R², R³, R⁴, R⁵, R⁶ and R⁷ are hydrogen and R⁸ is methyl.

A different procedure is preferred for preparing compounds of Formula I in which R⁴ is lower alkyl and R¹-R³ and R⁵-R⁸ are hydrogen. An example where R⁴ is methyl is shown in Reaction Scheme IB.

REACTION SCHEME IB



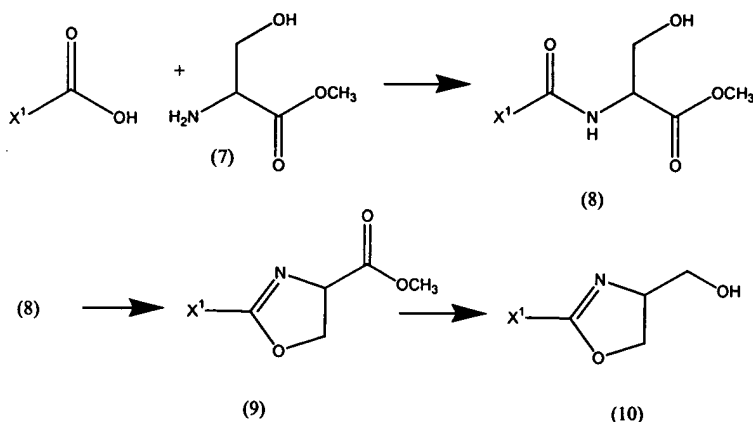
Step 1 - Preparation of Formula (7a)

The compound of formula (7) is reacted with 2-methylpiperazine. In general, the two compounds are mixed in an inert solvent, preferably a protic solvent, for example ethanol, in the presence of an inorganic or tertiary organic base, preferably triethylamine. The reaction is conducted at about 30-100°C, preferably at about 60°C, for about 2-12 hours, preferably about 8 hours. When the reaction is substantially complete, the product of formula (7a) is isolated by conventional means, for example by removal of the solvent under reduced pressure, followed by chromatography.

The compound of formula (7a) is then reacted with an epoxide of formula (3) as described in Reaction Scheme I, step 2, to provide a compound of Formula I in which R^1 , R^2 , R^3 , R^5 , R^6 , R^7 and R^8 are hydrogen and R^4 is methyl.

A preparation of intermediates for synthesizing compounds of Formula I in which Y is 2-substituted-(1,3-oxazolin-5-yl) is shown in Reaction Scheme II.

REACTION SCHEME II



Step 1 - Preparation of a Compound of Formula (8)

- 5 A carboxylic acid of formula $X^1C(O)OH$, is reacted with commercially available methyl 2-amino-3-hydroxypropanoate (7). In general, the two compounds are mixed in an inert solvent, for example dichloromethane, in the presence of an inorganic or tertiary organic base, preferably triethylamine. The reaction is conducted at about room temperature overnight. When the reaction is substantially complete, the product of
- 10 formula (8) is isolated by conventional means, for example by removal of the solvent under reduced pressure, followed by chromatography.

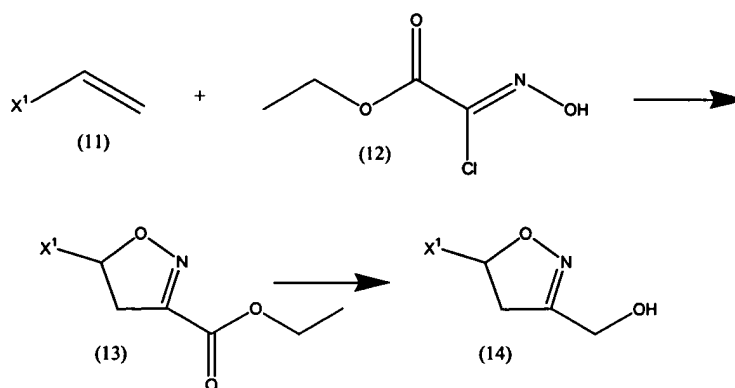
Step 2 - Preparation of a Compound of Formula (9)

- The compound of formula (8) is then cyclized by reaction with
- 15 diisopropylazodicarboxylate, or the like, in the presence of triphenylphosphine, to provide a 4-carbomethoxy-1,3-oxazoline of formula (9). The reaction is conducted in an inert solvent, for example tetrahydrofuran, at about room temperature for 1-5 days. When the reaction is substantially complete, the product of formula (9) is isolated by conventional means, for example by removal of the solvent under reduced pressure,
- 20 followed by chromatography.

Step 3 - Preparation of a Compound of Formula (10)

The carbomethoxy group of the compound of formula (9) is then reduced by conventional means to a hydroxymethyl group, to provide a compound of formula (10), for example by reduction with lithium aluminum hydride in an ethereal solvent, for

REACTION SCHEME III



Step 1 - Preparation of a Compound of Formula (13)

5 A vinyl derivative of formula (11) is reacted with ethyl 2-chloro-2-(hydroxyamino)acetate (12) in an inert solvent, for example tetrahydrofuran, in the presence of a tertiary base, for example triethylamine, for about 30 minutes to 4 hours. When the reaction is substantially complete, the product of formula (13) is isolated by conventional means.

10

Step 2 - Preparation of a Compound of Formula (14)

The carboxyethyl group of the compound of formula (13) is then reduced by conventional means to a hydroxymethyl group, to provide a compound of formula (14), for example by reduction with sodium borohydride in a protic solvent, for example ethanol, at about 0°C, for about 2-8 hours. When the reaction is substantially complete, the product of formula (14) is isolated by conventional means, for example by quenching excess reducing agent with water, extraction with an inert solvent, for example ethyl acetate, removal of the solvent under reduced pressure, followed by chromatography.

15

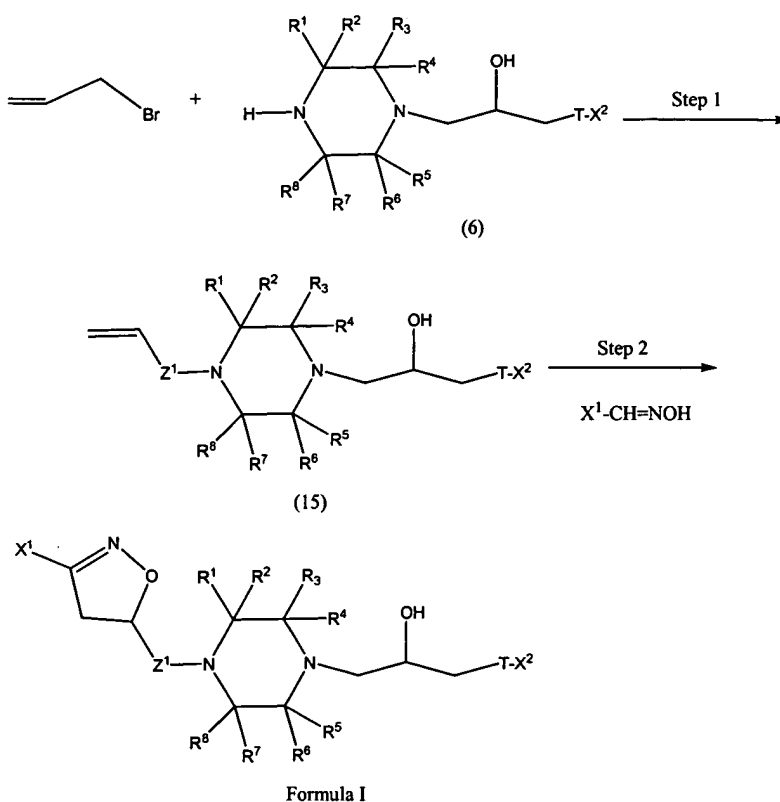
The hydroxymethyl of formula (14) may then be converted to a chloromethyl group by conventional means, for example thionyl chloride. or alternatively the hydroxy group may be converted to a mesyl group (leaving group) by reaction with methanesulfonyl chloride. The compound thus formed is then reacted with a piperazine derivative of formula (6) in the same manner as shown in Reaction Scheme I to provide a compound of Formula I in which Y is 5-substituted-(4,5-dihydroisoxazol-3-yl).

20

A method for preparing compounds of Formula I in which Y is 3-substituted-(4,5-dihydroisoxazol-3-yl) is shown in Reaction Scheme IV.

5

REACTION SCHEME IV



Step 1 - Preparation of a Compound of Formula (15)

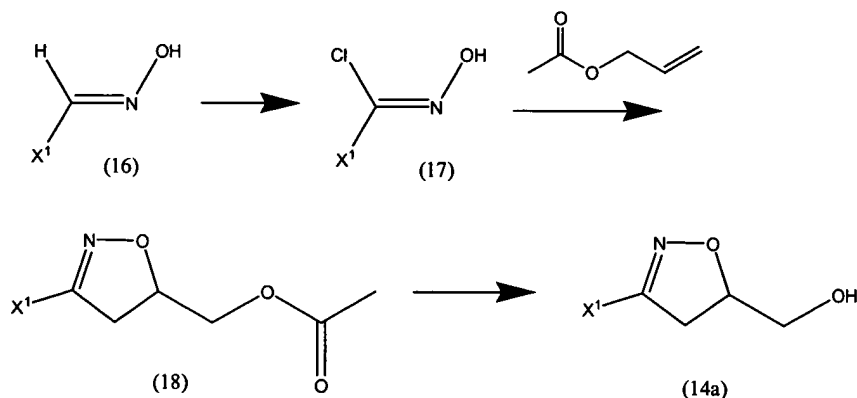
- 10 A compound of formula (6) is reacted with a compound of formula
 HC=C-Z¹Br (for example, allyl bromide). In general, the two compounds are mixed in
 an inert solvent, for example acetone, in the presence of an inorganic or tertiary organic
 base, preferably potassium carbonate. The reaction is conducted at about 30-100°C,
 preferably at about reflux, for about 1-10 hours, preferably about 5 hours. When the
 15 reaction is substantially complete, the product of formula (15) is isolated and purified by
 conventional means, for example by removal of the solvent under reduced pressure,
 followed by preparative chromatography.

Step 2 - Preparation of a Compound of Formula I

The compound of formula (15) is then reacted with a nitrile oxide, prepared by oxidation of an oxime of formula $X^1-CH=NOH$. The oxidation is carried out conventionally, for example by treatment of the oxime with an oxidizing agent such as aqueous sodium hypochlorite in dichloromethane. To the nitrile oxide thus prepared, the compound of formula (15) is added in an inert solvent, for example dichloromethane, in the presence of an inorganic or tertiary organic base, preferably triethylamine. The reaction is carried out in a sealed vessel at an elevated temperature, preferably subjecting the sealed vessel to microwave at about 110°C for about 4 minutes. When the reaction is substantially complete, the product of Formula I is isolated and purified by conventional means, for example by removal of the solvent under reduced pressure, followed by preparative chromatography. This provides a compound of Formula I in which Y is 3-substituted-(4,5-dihydroisoxazol-5-yl).

An alternative method for preparing compounds of Formula I in which Y is 3-substituted-(4,5-dihydroisoxazol-5-yl) is shown in Reaction Scheme V.

REACTION SCHEME V



Step 1 - Preparation of a Compound of Formula (17)

An oxime of formula (16), which is commercially available or prepared by means well known in the art, is reacted with a halogenating agent, for example N-chlorosuccinimide. In general, the two compounds are mixed in an inert solvent, for example dichloromethane, and reacted at about 0-30°C, preferably at about room

temperature, for about 1-10 hours, preferably about 3 hours. When the reaction is substantially complete, the product of formula (17) is typically used in the next reaction with no purification.

5 Step 2 - Preparation of a Compound of Formula (18)

 The compound of formula (17) is then reacted with a vinylic ester, for example vinyl acetate. The reaction is carried out in an inert solvent, for example dichloromethane, in the presence of an inorganic or tertiary organic base, for example triethylamine. The reaction is carried out at about 0-30°C, preferably at about room
10 temperature, for about 10-48 hours, preferably overnight. When the reaction is substantially complete, the product of formula (18) is isolated and purified by conventional means, for example by removal of the solvent under reduced pressure, followed by preparative chromatography.

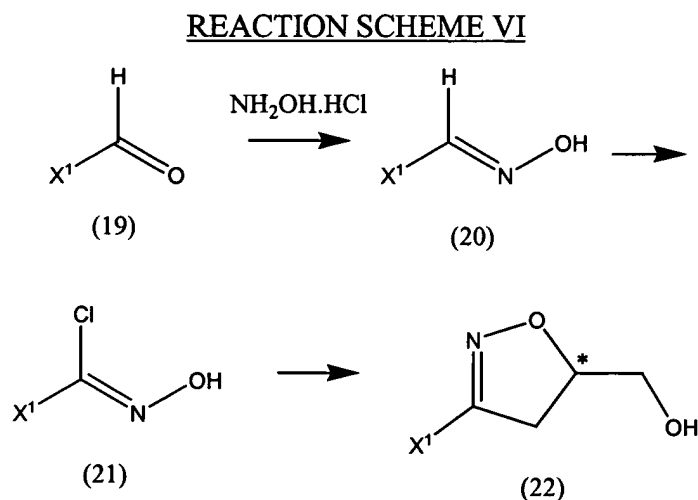
15 Step 3 - Preparation of a Compound of Formula (14a)

 The compound of formula (18) is then hydrolyzed to the hydroxymethyl compound of formula (14a) by reaction with a strong base, for example sodium hydroxide. The reaction is carried out in a protic solvent, for example ethanol-water mixture, at a temperature of about 0-30°C, preferably at about room temperature, for
20 about 1-6 hours, preferably for about 3 hours. When the reaction is substantially complete, the resulting solution is neutralized, and the product of formula (14a) is isolated and purified by conventional means.

 The hydroxy group of the compound of formula (14a) is then converted to a leaving group, for example a chloro or mesyl group, by conventional means, as shown
25 previously. The resulting halomethyl or mesylmethyl compound is then reacted with a piperazine derivative of formula (6) in the same manner as shown in Reaction Scheme I to provide a compound of Formula I in which Y is 3-substituted-(4,5-dihydroisoxazol-5-yl).

An example of a method for preparing compounds of Formula I in which X¹ is optionally substituted aryl or optionally substituted heteroaryl and Y is optically active is shown in Reaction Scheme VI.

5



Step 1 – Preparation of Compound of Formula (20)

- 10 The compound of formula (20) is prepared by reacting a compound of formula (19) with hydroxylamine hydrochloride (NH₂OH.HCl). In general, the two compounds are mixed in a protic solvent, for example methanol, in the presence of an inorganic or tertiary organic base, preferably potassium carbonate. The reaction is conducted at about 30-100°C, preferably at about 60-70°C, for about 2-20 hours, preferably about 10 hours.
- 15 When the reaction is substantially complete, the product of formula (20) is isolated and purified by conventional means, for example by filtration and removal of solvent under reduced pressure, followed by preparative chromatography.

Step 2 – Preparation of Compound of Formula (21)

- 20 The compound of formula (20) is reacted with a halogenating agent, preferably N-chlorosuccinimide (NCS) in a polar inert solvent, preferably dimethylformamide. In general, the reaction is conducted initially at a low temperature, preferably about 0°C, and then at about 30-70°C for about 1-10 hours, preferably 3 hours. When the reaction is substantially complete, the product of formula (21) is isolated and purified by

conventional means, for example by removal of solvent under reduced pressure, followed by preparative chromatography of the residue.

Step 3 – Preparation of Compound of Formula (22)

5 A solution of allyl alcohol in an aprotic solvent is reacted with diethylzinc at a low temperature, preferably about 0°C. The resulting mixture is then reacted with a chiral tartrate, for example (+)diisopropyl tartrate (+) or (-)diisopropyl tartrate for 1-10 hours, preferably 2 hours, after which another portion of diethylzinc is added. The compound of formula (21) is then added to the mixture thus produced, and reacted at 0°C
10 for another 1-10 hours, preferably about 2 hours. When the reaction is substantially complete, the product of formula (22) is isolated and purified by conventional means, for example by conventional workup and removal of solvent under reduced pressure, followed by preparative chromatography of the residue.

 The hydroxy group of the compound of formula (22) is then converted to a
15 leaving group, for example a chloro or mesyl group, by conventional means, as shown previously. The resulting halomethyl or mesylmethyl compound is then reacted with a piperazine derivative of formula (6) in the same manner as shown in Reaction Scheme I to provide a compound of Formula I in which X¹ is optionally substituted aryl or optionally substituted heteroaryl and Y is optically active.

General Utility

 The compounds of Formula I are effective in the treatment of conditions known to respond to administration of fatty acid oxidation inhibitors, including protection of skeletal muscles against damage resulting from trauma, intermittent claudication, shock,
25 and cardiovascular diseases including atrial and ventricular arrhythmias, Prinzmetal's (variant) angina, stable angina, ischemia and reperfusion injury in cardiac, kidney, liver and the brain, exercise induced angina, congestive heart disease, and myocardial infarction. Fatty acid oxidation inhibitors have recently been shown to modify glucose levels in diabetic patients, thus providing a novel method of treating diabetes, and in
30 particular provide an effective treatment of angina in diabetics. Fatty acid oxidation inhibitors have also been shown to raise plasma HDL levels and lower LDL levels in

mammals, thus providing a method for treating coronary artery disease. The compounds of Formula I can also be used to preserve donor tissue and organs used in transplants, and may be coadministered with thrombolytics, anticoagulants, and other agents.

Testing

5 Activity testing is conducted as described in those patents and patent applications referenced above, and in the Examples below, and by methods apparent to one skilled in the art.

Pharmaceutical Compositions

10 The compounds of Formula I are usually administered in the form of pharmaceutical compositions. This invention therefore provides pharmaceutical compositions that contain, as the active ingredient, one or more of the compounds of Formula I, or a pharmaceutically acceptable salt or ester thereof, and one or more pharmaceutically acceptable excipients, carriers, including inert solid diluents and fillers, diluents, including sterile aqueous solution and various organic solvents, permeation
15 enhancers, solubilizers and adjuvants. The compounds of Formula I may be administered alone or in combination with other therapeutic agents. Such compositions are prepared in a manner well known in the pharmaceutical art (see, e.g., Remington's Pharmaceutical Sciences, Mace Publishing Co., Philadelphia, PA 17th Ed. (1985) and "Modern
Pharmaceutics", Marcel Dekker, Inc. 3rd Ed. (G.S. Banker & C.T. Rhodes, Eds.).

20 Administration

 The compounds of Formula I may be administered in either single or multiple doses by any of the accepted modes of administration of agents having similar utilities, for example as described in those patents and patent applications incorporated by reference, including rectal, buccal, intranasal and transdermal routes, by intra-arterial
25 injection, intravenously, intraperitoneally, parenterally, intramuscularly, subcutaneously, orally, topically, as an inhalant, or via an impregnated or coated device such as a stent, for example, or an artery-inserted cylindrical polymer.

 One mode for administration is parental, particularly by injection. The forms in which the novel compositions of the present invention may be incorporated for
30 administration by injection include aqueous or oil suspensions, or emulsions, with sesame oil, corn oil, cottonseed oil, or peanut oil, as well as elixirs, mannitol, dextrose, or a

sterile aqueous solution, and similar pharmaceutical vehicles. Aqueous solutions in saline are also conventionally used for injection, but less preferred in the context of the present invention. Ethanol, glycerol, propylene glycol, liquid polyethylene glycol, and the like (and suitable mixtures thereof), cyclodextrin derivatives, and vegetable oils may also be employed. The proper fluidity can be maintained, for example, by the use of a coating, such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. The prevention of the action of microorganisms can be brought about by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the like.

Sterile injectable solutions are prepared by incorporating the compound of Formula I in the required amount in the appropriate solvent with various other ingredients as enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the various sterilized active ingredients into a sterile vehicle which contains the basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the preferred methods of preparation are vacuum-drying and freeze-drying techniques which yield a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

Compounds of Formula I may be impregnated into a stent by diffusion, for example, or coated onto the stent such as in a gel form, for example, using procedures known to one of skill in the art in light of the present disclosure.

Oral administration is another route for administration of the compounds of Formula I. Administration may be via capsule or enteric coated tablets, or the like. In making the pharmaceutical compositions that include at least one compound of Formula I, the active ingredient is usually diluted by an excipient and/or enclosed within such a carrier that can be in the form of a capsule, sachet, paper or other container. When the excipient serves as a diluent, it can be in the form of a solid, semi-solid, or liquid material (as above), which acts as a vehicle, carrier or medium for the active ingredient. Thus, the compositions can be in the form of tablets, pills, powders, lozenges, sachets, cachets, elixirs, suspensions, emulsions, solutions, syrups, aerosols (as a solid or in a liquid

medium), ointments containing, for example, up to 10% by weight of the active compound, soft and hard gelatin capsules, sterile injectable solutions, and sterile packaged powders.

Some examples of suitable excipients include lactose, dextrose, sucrose, sorbitol, mannitol, starches, gum acacia, calcium phosphate, alginates, tragacanth, gelatin, calcium silicate, microcrystalline cellulose, polyvinylpyrrolidone, cellulose, sterile water, syrup, and methyl cellulose. The formulations can additionally include: lubricating agents such as talc, magnesium stearate, and mineral oil; wetting agents; emulsifying and suspending agents; preserving agents such as methyl- and propylhydroxy-benzoates; sweetening agents; and flavoring agents.

The compositions of the invention can be formulated so as to provide quick, sustained or delayed release of the active ingredient after administration to the patient by employing procedures known in the art. Controlled release drug delivery systems for oral administration include osmotic pump systems and dissolutional systems containing polymer-coated reservoirs or drug-polymer matrix formulations. Examples of controlled release systems are given in U.S. Patent Nos. 3,845,770; 4,326,525; 4,902,514; and 5,616,345. Another formulation for use in the methods of the present invention employs transdermal delivery devices ("patches"). Such transdermal patches may be used to provide continuous or discontinuous infusion of the compounds of the present invention in controlled amounts. The construction and use of transdermal patches for the delivery of pharmaceutical agents is well known in the art. *See, e.g.*, U.S. Patent Nos. 5,023,252, 4,992,445 and 5,001,139. Such patches may be constructed for continuous, pulsatile, or on demand delivery of pharmaceutical agents.

The compositions are preferably formulated in a unit dosage form. The term "unit dosage forms" refers to physically discrete units suitable as unitary dosages for human subjects and other mammals, each unit containing a predetermined quantity of active material calculated to produce the desired therapeutic effect, in association with a suitable pharmaceutical excipient (e.g., a tablet, capsule, ampoule). The compounds of Formula I are effective over a wide dosage range and are generally administered in a pharmaceutically effective amount. Preferably, for oral administration, each dosage unit contains from 1 mg to 2 g of a compound of Formula I, and for parenteral administration,

preferably from 0.1 to 700 mg of a compound of Formula I. It will be understood, however, that the amount of the compound of Formula I actually administered will be determined by a physician, in the light of the relevant circumstances, including the condition to be treated, the chosen route of administration, the actual compound
5 administered and its relative activity, the age, weight, and response of the individual patient, the severity of the patient's symptoms, and the like.

For preparing solid compositions such as tablets, the principal active ingredient is mixed with a pharmaceutical excipient to form a solid preformulation composition containing a homogeneous mixture of a compound of the present invention. When
10 referring to these preformulation compositions as homogeneous, it is meant that the active ingredient is dispersed evenly throughout the composition so that the composition may be readily subdivided into equally effective unit dosage forms such as tablets, pills and capsules.

The tablets or pills of the present invention may be coated or otherwise
15 compounded to provide a dosage form affording the advantage of prolonged action, or to protect from the acid conditions of the stomach. For example, the tablet or pill can comprise an inner dosage and an outer dosage component, the latter being in the form of an envelope over the former. The two components can be separated by an enteric layer that serves to resist disintegration in the stomach and permit the inner component to pass
20 intact into the duodenum or to be delayed in release. A variety of materials can be used for such enteric layers or coatings, such materials including a number of polymeric acids and mixtures of polymeric acids with such materials as shellac, cetyl alcohol, and cellulose acetate.

Compositions for inhalation or insufflation include solutions and suspensions in
25 pharmaceutically acceptable, aqueous or organic solvents, or mixtures thereof, and powders. The liquid or solid compositions may contain suitable pharmaceutically acceptable excipients as described *supra*. Preferably the compositions are administered by the oral or nasal respiratory route for local or systemic effect. Compositions in preferably pharmaceutically acceptable solvents may be nebulized by use of inert gases.
30 Nebulized solutions may be inhaled directly from the nebulizing device or the nebulizing device may be attached to a face mask tent, or intermittent positive pressure breathing

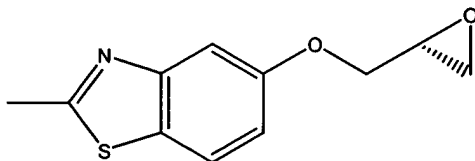
machine. Solution, suspension, or powder compositions may be administered, preferably orally or nasally, from devices that deliver the formulation in an appropriate manner.

The following examples are included to demonstrate preferred embodiments of the invention. It should be appreciated by those of skill in the art that the techniques disclosed in the examples which follow represent techniques discovered by the inventor to function well in the practice of the invention, and thus can be considered to constitute preferred modes for its practice. However, those of skill in the art should, in light of the present disclosure, appreciate that many changes can be made in the specific embodiments which are disclosed and still obtain a like or similar result without departing from the spirit and scope of the invention.

EXAMPLE 1

Preparation of a Compound of Formula (3)

A. Preparation of a Compound of Formula (3) in which T is Oxygen, X² is 2-Methylbenzothiazol-5-yl, and Z² is Methylene



A mixture of 2-methylbenzothiazol-5-ol, a compound of formula (1) (6.0 g, 36 mmol), (S)-(+)-epichlorohydrin, a compound of formula (2) (20 ml, 182 mmol), and potassium carbonate (20 g, 144 mmol) in acetone (100 ml), was heated to reflux and allowed to stir overnight. The solution was allowed to cool and filtered through Celite 512. The filtrate was evaporated (in vacuo), to yield an oil. The oil was chromatographed on silica gel, eluting with 20% ethyl acetate/hexanes, to yield 5-(((2R)oxiran-2-yl)methoxy)-2-methylbenzothiazole as a white solid (6.2 g, 28 mmol).

B. Preparation of a Compound of Formula (3) in which T is Oxygen, X² is 2-Phenylbenzothiazol-5-yl, and Z² is Methylene

Similarly, following the procedure of 1A above, but replacing 2-methylbenzothiazol-5-ol with 2-phenylbenzoxazol-5-ol, a compound of formula (3) where X² is 2-phenylbenzoxazol-5-yl, T is oxygen, and Z² is methylene was prepared, namely 5-(((2R)oxiran-2-yl)methoxy)-2-phenylbenzoxazole.

Similarly, the following compounds of formula (3) were prepared:

4-(((2R)oxiran-2-yl)methoxy)-1-methoxybenzene; and

4-(((2R)oxiran-2-yl)methoxy)-2-fluorobenzene.

C. Preparation of a Compound of Formula (3), varying T, X², and Z²

Similarly, following the procedure of 1A above, but optionally replacing 2-methylbenzothiazol-5-ol with other compounds of formula (1), and optionally replacing (S)-(+)-epichlorohydrin with other appropriately substituted compounds of formula (2), the following compounds of formula (3) are prepared as R, S isomers, or racemic mixtures.

5-(((2S)oxiran-2-yl)methoxy)-2-methylbenzothiazole;

5-(((2RS)oxiran-2-yl)methoxy)-2-methylbenzothiazole

1-[2-((2S)oxiran-2-yl)ethoxy]-2-methoxybenzene

4-(((2R)oxiran-2-yl)methoxy)-2-chlorobenzene;

5-[2-((2S)oxiran-2-yl)ethoxy]-2-methylbenzothiazole

4-[(oxiran-2-yl)ethoxy]-2-fluorobenzene;

4-[(oxiran-2-yl)ethoxy]-4-methoxybenzene;

8-fluoro-1-(oxiran-2-ylmethoxy)naphthalene;

1-fluoro-2-(oxiran-2-ylmethoxy)naphthalene;

2-ethyl-4-(oxiran-2-yl methoxy)thiazole;

4-methyl-2-(oxiran-2-yl methoxy)imidazole;

2-methyl-5-(oxiran-2-yl methoxy)benzimidazole; and

2-phenyl-5-(oxiran-2-yl methoxy)benzimidazole.

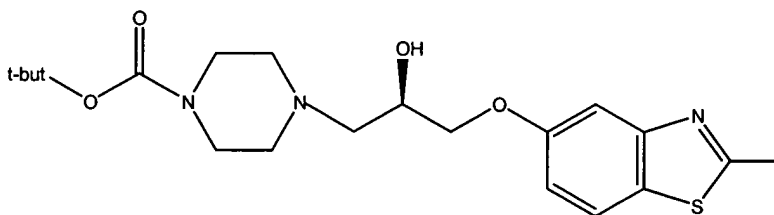
D. Preparation of a Compound of Formula (3), varying T, X², and Z²

Similarly, following the procedure of 1A above, but optionally replacing 2-methylbenzothiazol-5-ol with other compounds of formula (1), and optionally replacing (S)-(+)-epichlorohydrin with other appropriately substituted compounds of formula (2),
5 other compounds of formula (3) are prepared.

EXAMPLE 2

Preparation of a Compound of Formula (5)

10 A. Preparation of a Compound of Formula (5) in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are Hydrogen, T is Oxygen, X² is 2-Methylbenzothiazol-5-yl, and Z² is Methylene



To a solution of 5-[(2R)oxiran-2-yl)methoxy]-2-methylbenzothiazole, a
15 compound of formula (3) (6.2 g, 28 mmol), and *tert*-butyl 1-piperazinecarboxylate, a compound of formula (4) (5.7 g, 31 mmol), in methylene chloride (200 ml), was added ytterbium (III) trifluoromethanesulfonate (1.73g, 28 mmol). The resulting solution was allowed to stir at room temperature overnight. The solvent was evaporated (in vacuo), to yield a semi-solid, which was chromatographed on silica gel, eluting with 5%
20 methanol/methylene chloride, to yield *tert*-butyl 4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl]piperazinecarboxylate as a clear oil (9.5 g, 23 mmol).

B. Preparation of a Compound of Formula (5) in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are Hydrogen, T is Oxygen, X² is 2-Phenylbenzoxazol-5-yl, and Z² is Methylene

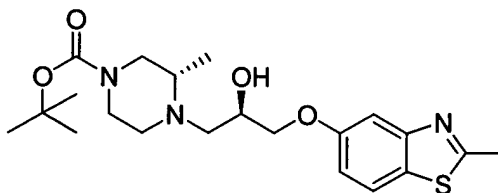
25 Similarly, following the procedure of 2A above, but replacing 5-[(2R)oxiran-2-yl)methoxy]-2-methylbenzothiazole with other compounds of formula (3), the following compounds of formula (5) were prepared:

tert-butyl 4-[(2R)-2-hydroxy-3-(2-phenylbenzoxazol-5-yloxy)propyl]-
piperazinecarboxylate;

tert-butyl 4-[(2R)-2-hydroxy-3-(2-methoxyphenoxy)propyl]piperazinecarboxylate; and
tert-butyl 4-[(2R)-2-hydroxy-3-(2-fluorophenoxy)propyl]piperazinecarboxylate.

5

C. Alternative Preparation of a Compound of Formula (5) in which R¹, R², R⁴, R⁵,
R⁶, R⁷, and R⁸ are Hydrogen, R³ is Methyl, T is Oxygen, X² is 2-Phenylbenzoxazol-5-yl,
and Z² is Methylene



10

To a solution of 5-(((R)-oxiran-2-yl)methoxy)-2-methylbenzo[d]thiazole (3.24 g,
14.6 mmol) in *tert*-amyl alcohol (50 ml) was added (*S*)-*tert*-butyl 3-methylpiperazine-1-
carboxylate (2.67 g, 13.3 mmol). The resulting mixture was stirred at 100°C for 16 hours.
It was then cooled, concentrated, and purified by chromatography (1:20 MeOH:CH₂Cl₂)
15 to yield tert-butyl 4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl](3S)-3-
methylpiperazinecarboxylate (4.8 g, 11.4 mmol).

D. Preparation of a Compound of Formula (5), varying R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸,
T, X², and Z²

20

Similarly, following the procedure of 2A or 2C above, but optionally replacing 5-
[(((2R)oxiran-2-yl)methoxy)]-2-methylbenzothiazole with other compounds of formula
(3), and optionally replacing *tert*-butyl 1-piperazinecarboxylate with other compounds of
formula (4), the following compounds of formula (5) are prepared as racemates or single
enantiomers:

25

tert-butyl 4-[2-hydroxy-3-(2-methoxyphenoxy)propyl]piperazinecarboxylate;
tert-butyl 4-[2-hydroxy-3-(4-methoxyphenoxy)propyl]piperazinecarboxylate;
tert-butyl 4-[2-hydroxy-3-(2-fluorophenoxy)propyl]piperazinecarboxylate;
tert-butyl 4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl]-
piperazinecarboxylate;

4-[2-hydroxy-3-(8-fluoronaphth-1-yloxy)propyl]-piperazine-1-carboxylic acid *tert*-butyl ester;

4-[2-hydroxy-3-(1-fluoronaphth-2-yloxy)propyl]-piperazine-1-carboxylic acid *tert*-butyl ester;

5 4-[2-hydroxy-3-(2-ethylthiazol-4-yloxy)propyl]-piperazine-1-carboxylic acid *tert*-butyl ester;

4-[2-hydroxy-3-(4-methylimidazol-4-yloxy)propyl]-piperazine-1-carboxylic acid *tert*-butyl ester;

10 4-[2-hydroxy-3-(2-methylbenzimidazol-5-yloxy)propyl]-piperazine-1-carboxylic acid *tert*-butyl ester; and

4-[2-hydroxy-3-(2-phenylbenzimidazo-5-yloxy)propyl]-piperazine-1-carboxylic acid *tert*-butyl ester.

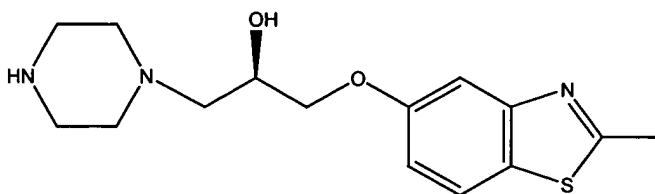
15 E. Preparation of a Compound of Formula (5), varying R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, T, X², and Z²

Similarly, following the procedure of 2A or 2C above, but optionally replacing 5-(((2R)oxiran-2-yl)methoxy)-2-methylbenzothiazole with other compounds of formula (3), and optionally replacing *tert*-butyl 1-piperazinecarboxylate with other compounds of formula (4), other compounds of formula (5) are prepared as racemates or single enantiomers.

EXAMPLE 3

Preparation of a Compound of Formula (6)

A. Preparation of a Compound of Formula (6) in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are Hydrogen, T is Oxygen, X² is 2-Methylbenzothiazol-5-yl, and Z² is Methylene



A solution of tert-butyl 4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl]piperazinecarboxylate, a compound of formula (5) (2.0 g, 4.9 mmol), and 25% trifluoroacetic acid /methylene chloride (20 ml) was allowed to stir at room temperature overnight. The solvent was evaporated (in vacuo) to yield an oil. The oil was diluted with acetone (20 ml) and solid potassium carbonate was added until the foaming stopped. The resulting mixture was allowed to stir overnight. The solution was filtered through Celite 512, and the filtrate was evaporated (in vacuo), to yield an oil. The oil was placed under high vacuum overnight, to yield (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazinylpropan-2-ol as a clear viscous oil (3.4 g, 6.3 mmol).

B. Preparation of a Compound of Formula (6) in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are Hydrogen, T is Oxygen, X² is 2-Phenylbenzoxazol-5-yl, and Z² is Methylene

Similarly, following the procedure of 3A above, but replacing tert-butyl 4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl]piperazinecarboxylate with tert-butyl 4-[(2R)-2-hydroxy-3-(2-phenylbenzoxazol-5-yloxy)propyl]piperazinecarboxylate, the compound of formula (6) where R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are hydrogen, X² is 2-phenylbenzoxazol-5-yl, T is oxygen, and Z² is methylene was prepared, namely (2R)-1-(2-phenylbenzoxazol-5-yloxy)-3-piperazinylpropan-2-ol.

Similarly, the following compounds of formula (6) were prepared:

(2R)-1-((2S)-2-methylpiperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;
(2R)-1-(2-methoxyphenoxy)-3-piperazinylpropan-2-ol; and
(2R)-1-(2-fluorophenoxy)-3-piperazinylpropan-2-ol.

C. Preparation of a Compound of Formula (6), varying R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, T, X², and Z²

Similarly, following the procedure of 3A above, but replacing tert-butyl 4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl]piperazinecarboxylate with other compounds of formula (5), the following compounds of formula (6) are prepared as racemates or single enantiomers:

- 4-(2-methoxyphenoxy)-1-piperazinylobutan-2-ol;
- 4-(2-chlorophenoxy)-1-piperazinylobutan-2-ol;
- 4-(2-methylbenzothiazol-5-yloxy)-1-piperazinylobutan-2-ol
- 4-(2-fluorophenoxy)-1-piperazinylopropan-2-ol;
- 4-(4-methoxyphenoxy)-1-piperazinylopropan-2-ol;
- 1-(8-fluoronaphthyl-2-yloxy)-3-piperazinylopropan-2-ol;
- 1-(1-fluoronaphth-2-yloxy)-3-piperazinylopropan-2-ol;
- 1-(2-ethylthiazol-4-yloxy)-3-piperazinylopropan-2-ol;
- 1-(4-methylimidazol-4-yloxy)-3-piperazinylopropan-2-ol;
- 1-(2-methylbenzimidazol-5-yloxy)-3-piperazinylopropan-2-ol; and
- 1-(2-phenylbenzimidazol-5-yloxy)-3-piperazin-1-yl-propan-2-ol.

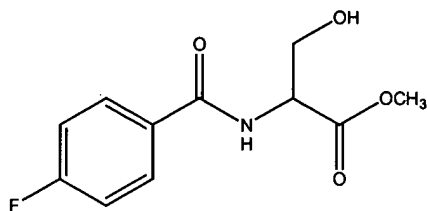
D. Preparation of a Compound of Formula (6), varying R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, T, X², and Z²

Similarly, following the procedure of 3A above, but replacing tert-butyl 4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl]piperazinecarboxylate with other compounds of formula (5), other compounds of formula (6) are prepared as racemates or single enantiomers.

EXAMPLE 4

Preparation of a Compound of Formula (8)

A. Preparation of a Compound of Formula (8) in which X¹ is 4-Fluorophenyl



5

To 50ml of tetrahydrofuran was added 4-fluorobenzoic acid (3.64g, 26mmol), L-serine methyl ester hydrochloride (2g, 13mmol), 1-hydroxybenzotriazole hydrate (4g, 26mmol), O-benzotriazol-1-yl-N,N,N',N'-tetramethyluronium hexafluorophosphate (9.9g, 26mmol), triethylamine (3.5ml, 26mmol) and dimethylaminopyridine (20mg, 0.16mmol). The reaction was stirred for 24 hours at room temperature. Solvent was removed under reduced pressure, and the residue dissolved in 150ml of ethyl acetate.

10

The organic solution was washed in a separatory funnel in the following manner: 3x100ml 10% aqueous citric acid, 3x100ml saturated sodium bicarbonate, 3x100ml water, 1x75ml saturate sodium chloride solution. The organic layer was concentrated and purified using a Flash40 column (1:2 EtOAc:Hexane, 20psi) to yield 3 grams of methyl 2-[(4-fluorophenyl)carbonylamino]-3-hydroxypropanoate.

15

B. Preparation of a Compound of Formula (8), varying X¹

Similarly, following the procedure of 4A above, but replacing 4-fluorobenzoic acid with other compounds of formula X¹-CO(O)H, other compounds of formula (8) are prepared as racemates or single enantiomers.

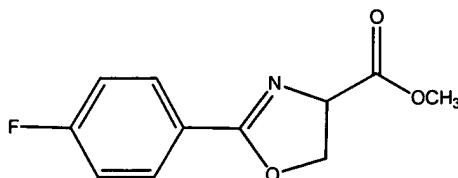
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25

EXAMPLE 5

Preparation of a Compound of Formula (9)

A. Preparation of a Compound of Formula (9) in which X¹ is 4-Fluorophenyl



5

To 25ml tetrahydrofuran was added methyl 2-[(4-fluorophenyl)carbonylamino]-3-hydroxypropanoate (0.750g, 3mmol), triphenylphosphine (1.0g, 4.5mmol) and diisopropylazodicarboxylate (0.800g, 4.5mmol). The reaction was stirred at room temperature under nitrogen until SM was shown to be consumed (TLC, 1:1 Ethyl
10 Acetate:Hexane). The reaction mixture was concentrated under reduced pressure and purified using a Flash 40 Column (1:1 EtOAc:Hexane, 20psi) to yield 1g of methyl 2-(4-fluorophenyl)-1,3-oxazoline-4-carboxylate (M+1 = 224.2).

B. Preparation of a Compound of Formula (9), varying X¹

15

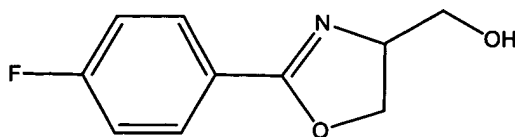
Similarly, following the procedure of 5A above, but replacing methyl 2-[(4-fluorophenyl)carbonylamino]-3-hydroxypropanoate with other compounds of formula (8), other compounds of formula (9) are prepared as racemates or single enantiomers.

20

EXAMPLE 6

Preparation of a Compound of Formula (10)

A. Preparation of a Compound of Formula (10) in which X¹ is 4-Fluorophenyl



25

Methyl 2-(4-fluorophenyl)-1,3-oxazoline-4-carboxylate (1.0g, 4.5mmol) was dissolved in 30ml of absolute ethanol. To this solution was added lithium chloride (0.380g, 9.0mmol) and sodium borohydride (0.342g, 9.0mmol). The reaction was stirred at room temperature until starting material was consumed (TLC, 15:1

dichloromethane:methanol). Ethanol was removed under reduced pressure and replaced with 35ml of water. The aqueous solution was acidified to about pH 2 with concentrated HCl, and the acidic solution extracted with ethyl acetate (3X50ml). The resulting organic layer was washed with water (3x50ml) and dried with sodium sulfate. The organic layer was concentrated under reduced pressure, and the residue purified using column chromatography (3% MeOH in DCM) to yield [2-(4-fluorophenyl)-1,3-oxazolin-4-yl]methan-1-ol (600mg, M+1 =196.1).

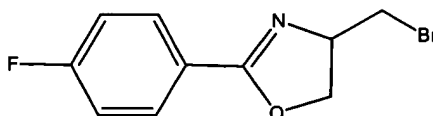
B. Preparation of a Compound of Formula (10), varying X¹

Similarly, following the procedure of 6A above, but replacing methyl 2-(4-fluorophenyl)-1,3-oxazoline-4-carboxylate with other compounds of formula (9), other compounds of formula (10) are prepared as racemates or single enantiomers.

EXAMPLE 7

Preparation of a Compound of Formula (10a)

A. Preparation of a Compound of Formula (10a) in which X¹ is 4-Fluorophenyl and LG is Bromo



To a solution of [2-(4-fluorophenyl)-1,3-oxazolin-4-yl]methan-1-ol (500mg, 2.5mmol) in pyridine (30ml) cooled to 0° under inert conditions was added triphenylphosphine (1.3g, 5.0mmol). Carbon tetrabromide (817mg, 2.5mmol) was added in 4 fractions (~200mg each). The reaction mixture was stirred at zero degrees until all starting material was consumed, as shown by TLC (1:1 EtOAc:Hexanes). Methanol (~5ml) was added to quench the reaction. The solution was evaporated under reduced pressure, and the residue was purified using column chromatography (1:1 EtOAc:Hexanes) to yield compound 4-(bromomethyl)-2-(4-fluorophenyl)-1,3-oxazoline (300mg, HNMR).

B. Preparation of a Compound of Formula (10a), varying X¹

Similarly, following the procedure of 7A above, but replacing [2-(4-fluorophenyl)-1,3-oxazolin-4-yl]methan-1-ol with other compounds of formula (10), other compounds of formula (10a) are prepared as racemates or single enantiomers.

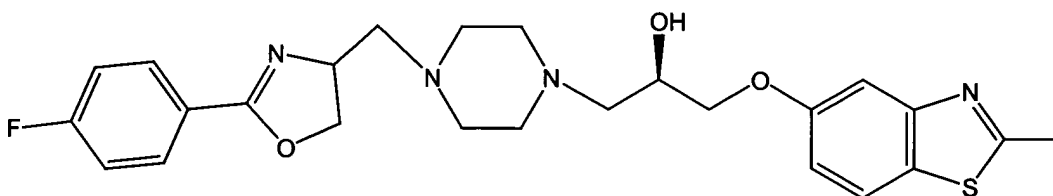
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EXAMPLE 8

Preparation of a Compound of Formula I

A. Preparation of a Compound of Formula I in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are Hydrogen, T is Oxygen, X¹-Y- is 2-(4-Fluorophenyl)-4,5-dihydro-1,3-oxazolin-4-yl, X² is 2-Methylbenzothiazol-5-yl, and Z¹ and Z² are Methylene

10



To a solution of 4-(bromomethyl)-2-(4-fluorophenyl)-1,3-oxazoline (80mg, 0.31mmol) in dimethylformamide (2ml) was added (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazinylpropan-2-ol (180mg, 0.47mmol) and potassium carbonate (250mg, 1.92mmol). The mixture was heated to 75 degrees and stirred for 12 hours. The reaction was then concentrated under reduced pressure and the residue purified using preparative chromatography (10:1 dichloromethane:methanol) to yield (2R)-3-(4-([2-(4-fluorophenyl)(4,5-dihydro-1,3-oxazolin-4-yl)methyl}piperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol (6mg, M+1 = 485.05).

15

20

B. Preparation of a Compound of Formula I, varying X¹

Similarly, following the procedure of 8A above, but optionally replacing 4-(bromomethyl)-2-(4-fluorophenyl)-4,5-dihydro-1,3-oxazoline with other compounds of formula (10a), and optionally replacing (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazine-1-yl-propan-2-ol with other compounds of formula (6), the following compounds of Formula I were prepared:

25

(2R)-1-{4-[[[(4S)-2-phenyl(1,3-oxazolin-4-yl)]methyl]piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-{4-[[[(4S,5S)-2-methyl-5-phenyl(1,3-oxazolin-4-yl)]methyl]piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol; and

5 (2R)-3-(4-{[2-(4-fluorophenyl)(1,3-oxazolin-4-yl)]methyl}piperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol.

C. Preparation of a Compound of Formula I, varying X¹

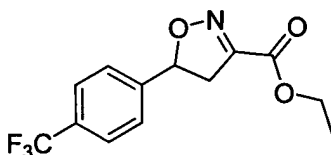
Similarly, following the procedure of 8A above, but optionally replacing 4-(bromomethyl)-2-(4-fluorophenyl)-4,5-dihydro-1,3-oxazoline with other compounds of
10 formula (10a), and optionally replacing (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazine-1-yl-propan-2-ol with other compounds of formula (6), other compounds of Formula I are prepared as racemates or single enantiomers.

15

EXAMPLE 9

Preparation of a Compound of Formula (13)

A. Preparation of a Compound of Formula (13) in which X¹ is 4-Trifluorophenyl



20

To a stirred mixture of 2-chloro-2-hydroxyiminoacetic acid ethyl ester (2.11g, 13.52mmol) and 4-trifluoromethylstyrene (2.0g, 11.27mmol) in anhydrous tetrahydrofuran (15ml) was added dropwise a solution of triethylamine (3.0ml, 21.52mmol) in tetrahydrofuran (5ml) at room temperature. The resulting reaction
25 mixture was stirred at an atmosphere of nitrogen for 16 hours, filtered, partitioned between water and ethyl acetate (2 x 50ml), washed with saturated aqueous sodium bicarbonate, 30% aqueous ammonium chloride, brine, dried over sodium sulfate, and concentrated to afford a pale yellow solid, which was purified by chromatography (2:8

EtOAc:Hexane) to afford ethyl 5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole-3-carboxylate (2.86g, M+1 = 288.2)

B. Preparation of a Compound of Formula (13), varying X¹

5 Similarly, following the procedure of 9A above, but replacing 4-trifluoromethystyrene with other compounds of formula (11), the following compounds of formula (13) were prepared:

ethyl 5-(2-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole-3-carboxylate;

ethyl 5-phenyl-4,5-dihydroisoxazole-3-carboxylate;

10 ethyl 5-(2,4,6-trimethylphenyl)-4,5-dihydroisoxazole-3-carboxylate;

ethyl 5-(2,6-dimethylphenyl)-4,5-dihydroisoxazole-3-carboxylate; and

ethyl 5-(2-chlorophenyl)-4,5-dihydroisoxazole-3-carboxylate.

C. Preparation of a Compound of Formula (13), varying X¹

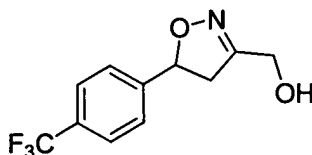
15 Similarly, following the procedure of 9A above, but replacing 4-trifluoromethystyrene with other compounds of formula (11), other compounds of formula (13) are prepared.

20

EXAMPLE 10

Preparation of a Compound of Formula (14)

A. Preparation of a Compound of Formula (14) in which X¹ is 4-Trifluorophenyl



25 To a stirred mixture of sodium borohydride (378mg, 10.0mmol) in ethanol (20ml) was added slowly a solution of ethyl 5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole-3-carboxylate (1.6g, 5.75mmol) in ethanol (10mL). The resulting reaction mixture was stirred under an atmosphere of nitrogen for 8 hours. Ethanol was removed under reduced pressure and water (20ml) added. The aqueous mixture was acidified to about pH 3 with concentrated HCl, and the acidic solution extracted with ethyl acetate (3X20ml). The

resulting organic phase was washed with saturated aqueous sodium bicarbonate (20ml), 30% aqueous ammonium chloride(20ml), brine, dried over sodium sulfate, concentrated, and purified using chromatography (3% MeOH in CH₂Cl₂) to afford (5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazol-3-yl)methanol (1.1g, M+1 = 246.1).

5

B. Preparation of a Compound of Formula (14), varying X¹

Similarly, following the procedure of 10A above, but replacing ethyl 5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole-3-carboxylate with other compounds of formula (13), the following compounds of formula (14) were prepared:

- 10 (5-(2-(trifluoromethyl)phenyl)-4,5-dihydroisoxazol-3-yl)methanol;
(5-phenyl-4,5-dihydroisoxazol-3-yl)methanol;
(5-(2,4,6-trimethylphenyl)-4,5-dihydroisoxazol-3-yl)methanol;
(5-(2,6-dimethylphenyl)-4,5-dihydroisoxazol-3-yl)methanol; and
(5-(2-chlorophenyl)-4,5-dihydroisoxazol-3-yl)methanol.

15

C. Preparation of a Compound of Formula (14), varying X¹

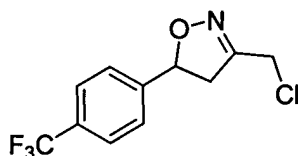
Similarly, following the procedure of 10A above, but replacing ethyl 5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole-3-carboxylate with other compounds of formula (13), other compounds of formula (14) are prepared.

20

EXAMPLE 11

Preparation of a Chloromethyl Derivative of a Compound of Formula (14)

A. Preparation of 3-(Chloromethyl)-5-[4-(trifluoromethyl)phenyl]-4,5-
25 dihydroisoxazole



To a stirred mixture of (5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazol-3-yl)methanol (245mg, 1.0mmol) in methylene chloride (15mL) was added thionyl chloride (200μL, 2.74mmol) dropwise at 0°C. The resulting mixture was stirred under an

atmosphere of nitrogen for 15 hours, concentrated under reduced pressure, and purified by preparative chromatography (3:7 EtOAc:hexane) to yield 3-(chloromethyl)-5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole (56mg, M+1 = 264.1).

5 B. Preparation of Similar Compounds

Similarly, following the procedure of 11A above, but replacing (5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazol-3-yl)methanol with other compounds of formula (14), the following chloromethyl derivatives were prepared:

3-(chloromethyl)-5-(2-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole;

10 3-(chloromethyl)-5-phenyl-4,5-dihydroisoxazole;

3-(chloromethyl)-5-(2,4,6-trimethylphenyl)-4,5-dihydroisoxazole;

3-(chloromethyl)-5-(2,6-dimethylphenyl)-4,5-dihydroisoxazole; and

3-(chloromethyl)-5-(2-chlorophenyl)-4,5-dihydroisoxazole.

15 C. Preparation of Similar Compounds

Similarly, following the procedure of 11A above, but replacing (5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazol-3-yl)methanol with other compounds of formula (14), other chloromethyl derivatives are prepared.

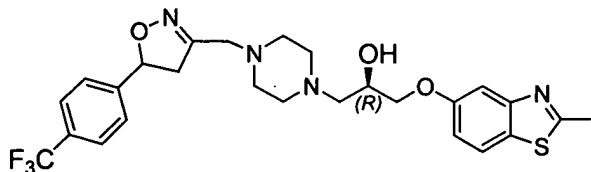
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EXAMPLE 12

Preparation of a Compound of Formula I

- A. Preparation of a Compound of Formula I in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are Hydrogen, T is Oxygen, X¹-Y- is 5-(4-trifluoromethylphenyl)-4,5-dihydroisoxazolin-3-yl, X² is 2-Methylbenzothiazol-5-yl, and Z¹ and Z² are Methylene



- To a stirred mixture of (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazinylpropan-2-ol (30mg, 0.08mmol) in *tert*-amylalcohol (5mL) was added triethylamine (40μL, 0.30mmol), followed by 3-(chloromethyl)-5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole (20mg, 0.08mmol). The resulting mixture was stirred at 95°C for 12 hours, concentrated under reduced pressure, and purified by preparative chromatograph (1:19 MeOH:CH₂Cl₂) to afford (2R)-3-(2-methylbenzothiazol-5-yloxy)-1-[4-((5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl))methyl)piperazinyl]propan-2-ol (10mg, M+1 = 535.1).

B. Preparation of a Compound of Formula I

- Similarly, following the procedure of 12A above, but optionally replacing (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazinylpropan-2-ol with other compounds of formula (6), and optionally replacing 3-(chloromethyl)-5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole with other chloromethyl derivatives, the following compounds of Formula I were prepared:

- (2R)-3-(2-methylbenzothiazol-5-yloxy)-1-{4-[(5-phenyl(4,5-dihydroisoxazol-3-yl))methyl]piperazinyl}propan-2-ol;
- (2R)-3-(2-methylbenzothiazol-5-yloxy)-1-(4-{[5-(2,4,6-trimethylphenyl)(4,5-dihydroisoxazol-3-yl)]methyl}piperazinyl)propan-2-ol;
- (2R)-3-(2-methylbenzothiazol-5-yloxy)-1-(4-{[5-(2,6-dimethylphenyl)(4,5-dihydroisoxazol-3-yl)]methyl}piperazinyl)propan-2-ol;

(2R)-3-(2-methylbenzothiazol-5-yloxy)-1-[4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-[(2S)-2-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

5 (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol; and

10 (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[2-chlorophenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol.

C. Preparation of a Compound of Formula I

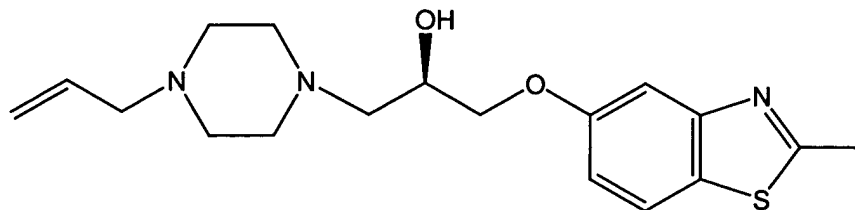
Similarly, following the procedure of 12A above, but optionally replacing (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazinylpropan-2-ol with other compounds of
15 formula (6), and optionally replacing 3-(chloromethyl)-5-(4-(trifluoromethyl)phenyl)-4,5-dihydroisoxazole with other chloromethyl derivatives, other compounds of Formula I are prepared.

20

EXAMPLE 13

Preparation of a Compound of Formula (15)

A. Preparation of a Compound of Formula (15) in which R¹, R², R³, R⁴, R⁵, R⁶, R⁷, and R⁸ are Hydrogen, T is Oxygen, X² is 2-Methylbenzothiazol-5-yl, and Z¹ and Z² are Methylene



25

To a solution of (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazin-1-yl-propan-2-ol (300mg, 1.0mmol) in acetone (10ml) was added allyl bromide (0.130ml, 1.5mmol)

and potassium carbonate (280mg, 2.0mmol). The solution was heated to reflux and stirred for 5 hours. Thin layer chromatography (10:1 dichloromethane:methanol) showed formation of product. The solution was cooled and filtered. The filtrate was concentrated under reduced pressure, and the residue purified by preparative chromatography (10:1 dichloromethane:methanol) to yield (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-(4-prop-2-enylpiperazinyl)propan-2-ol (150mg, M+1 = 348.2)

B. Preparation of a Compound of Formula (15)

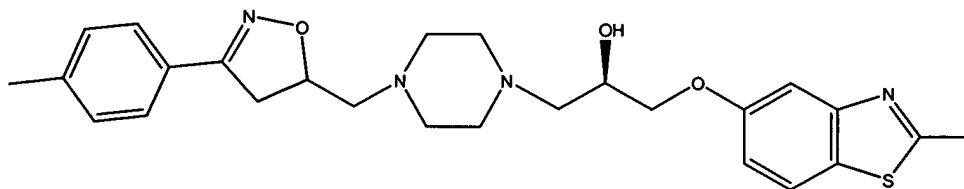
Similarly, following the procedure of 13A above, but optionally replacing (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-piperazin-1-yl-propan-2-ol with other compounds of formula (6), other compounds of formula (15) are prepared as racemates or single enantiomers.

EXAMPLE 14

Preparation of a Compound of Formula I

A. Preparation of a Compound of Formula I in which R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , and R^8 are Hydrogen, T is Oxygen, X^1 -Y- is 3-(4-Methylphenyl)-4,5-dihydroisoxazol-5-yl),

X^2 is 2-Methylbenzothiazol-5-yl, and Z^1 and Z^2 are Methylene



To a solution of (hydroxyimino)(4-methylphenyl)methane (97mg, 0.72mmol) in 1ml of dichloromethane was added aqueous sodium hypochlorite solution (>4%, 1.33ml, 0.72mmol). The reaction mixture was stirred at room temperature for 30 minutes, after which time TLC (20:1 dichloromethane:methanol) showed conversion of oxime to chlorinated oxime (17). 1-(2-Methylbenzothiazol-5-yloxy)-3-(4-prop-2-

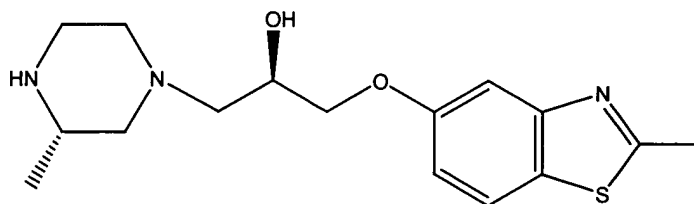
enylpiperazinyl)propan-2-ol (50mg, 0.144mmol) in 0.5ml of dichloromethane and triethylamine (0.022ml, 0.145mmol) were added.. The reaction was stirred at room temperature overnight. Thin layer chromatography (10:1 dichloromethane:methanol) showed consumption of starting material and product formation. The solution was
5 evaporated under reduced pressure, and the residue purified by preparative HPLC, to yield (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-(4-{[3-(4-methylphenyl)(4,5-dihydroisoxazol-5-yl)]methyl}piperazinyl)propan-2-ol (3.6mg, M+1 = 481.35)

Preparation of Starting Oxime

10 To a solution of 4-tolualdehyde (600mg, 5mmol) in tetrahydrofuran (3ml) was added hydroxylamine hydrochloride (700mg, 10mmol) and triethylamine (1.4ml, 10mmol). The reaction was subjected to microwave; absorbance normal; temp: 150°C; time; 600(s). The solid material was filtered off, and solvent removed from the filtrate under reduced pressure. The residue was purified using column chromatography (2:1
15 Hexanes:Ethyl Acetate) to provide (hydroxyimino)(4-methylphenyl)methane (500mg).

EXAMPLE 15

Preparation of a Compound of Formula (6) in which R¹, R², R³, R⁴, R⁵, R⁶ and R⁷ are
20 Hydrogen, R⁸ is (S)-Methyl, T is Oxygen, X² is 2-Methylbenzothiazol-5-yl, and Z² is Methylene



To a stirred solution of (2S)-methylpiperazine (1.0g, 10.0mmol) in ethanol (10mL) was added slowly a solution of 5-(((2R)oxiran-2-yl)methoxy)-2-methylbenzothiazole (2.20g, 10.0mmol) in ethanol (10mL). The resulting mixture was
25 stirred at 80°C under an atmosphere of nitrogen for 5 hours. It was then cooled, concentrated, and purified by chromatography (1:9 MeOH:CH₂Cl₂) to yield (2R)-3-((3S)-3-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol (2.28g, M+1 =

322.25), which was then treated with 4N HCl in 1,4-dioxane (4mL, 8.0mmol) followed by evaporation in vacuum to afford the hydrochloride salt as a pale yellow solid (2.80g, 7.6mmol).

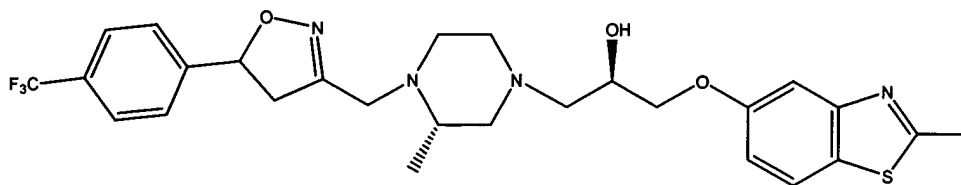
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EXAMPLE 16

Alternative Preparation of a Compound of Formula I

Preparation of a Compound of Formula I in which R¹, R², R³, R⁴, R⁵, R⁶ and R⁷ are Hydrogen, R⁸ is (S)-Methyl, T is Oxygen, X¹-Y- is 5-(4-Trifluorophenyl)-4,5-dihydroisoxazol-3-yl, X² is 2-Methylbenzothiazol-5-yl, and Z¹ and Z² are Methylene

10



To a stirred mixture of (2R)-3-((3S)-3-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol (32mg, 0.08mmol) in *tert*-amylalcohol (5mL) was added triethylamine (42μL, 0.30mmol), and 3-chloromethyl-5-(4-trifluorophenyl)isoxazole (20mg, 0.08mmol). The resulting mixture was stirred at 95°C for 12 hours. The reaction mixture was then cooled, concentrated under reduced pressure, and purified by preparative chromatograph (1:19 MeOH:CH₂Cl₂) to afford (2R)-3-[(3S)-3-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol (12mg, M+1 = 549.1).

15

20

B. Preparation of a Compound of Formula I

Similarly, following the procedure of 16A above, but optionally replacing (2R)-3-((3S)-3-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol with other compounds of formula (6), and optionally replacing 3-chloromethyl-5-(4-trifluorophenyl)isoxazole with other compounds of formula (7), the following compounds of Formula I were prepared:

25

(2R)-1-[4-({(5R)-3-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)}methyl)(2S)-2-methylpiperazinyl]-3-benzothiazol-5-yloxypropan-2-ol;

(2R)-1-{{(3S)-4-[[{(4S)-2-phenyl(1,3-oxazolin-4-yl)}methyl]-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

5 (2R)-1-{4-[[{(5R)-3-phenyl(4,5-dihydroisoxazol-5-yl)}methyl](2S)-2-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-{{(2S)-4-[[{(4S,5S)-2-methyl-5-phenyl(1,3-oxazolin-4-yl)}methyl]-2-methylpiperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[(2S)-2-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

10 (2R)-1-[(3S)-3-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-{4-[[{(5R)-3-phenyl(4,5-dihydroisoxazol-5-yl)}methyl](3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

15 N-{{[3-({4-[(2R)-2-hydroxy-3-(2-methylbenzothiazol-5-yloxy)propyl](2S)-2-methylpiperazinyl]carbonyl}(4,5-dihydroisoxazol-5-yl)]methyl}{2-chlorophenyl}carboxamide;

3-[(2S)-2-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl](2R)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

20 3-[(2S)-2-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl](2R)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-[(3S)-3-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-[(3S)-3-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

25 (2R)-3-((2S)-2-methyl-4-{{[3-(4-methylphenyl)(4,5-dihydroisoxazol-5-yl)]methyl}piperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

3-({[3-(4-chlorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl}(2S)-2-methylpiperazinyl)(2R)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

30 (2R)-1-{{(2S)-2-methyl-4-[(5-phenyl(4,5-dihydroisoxazol-3-yl)]methyl}piperazinyl}-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-((3S)-3-methyl-4-[(5-phenyl(4,5-dihydroisoxazol-3-yl)methyl]piperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-((3S)-4-[[5-(2,6-dimethylphenyl)(4,5-dihydroisoxazol-3-yl)]methyl]-3-methylpiperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

5 (2R)-1-(4-[[[(5R)-3-(4-fluorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl}(2S)-2-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-(4-[[[(5R)-3-(4-fluorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl}(3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

10 (2R)-1-[4-(([(5R)-3-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)(2S)-2-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-1-[4-(([(5R)-3-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)(3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol; and

(2R)-1-[4-(([(5R)-3-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl)]methyl)(3S)-3-methylpiperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol.

15

C. Preparation of a Compound of Formula I

Similarly, following the procedure of 16A above, but optionally replacing (2R)-3-((3S)-3-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol with other compounds of formula (6), and optionally replacing 3-chloromethyl-5-(4-trifluorophenyl)isoxazole with other compounds of formula (7), other compounds of Formula I are prepared.

20

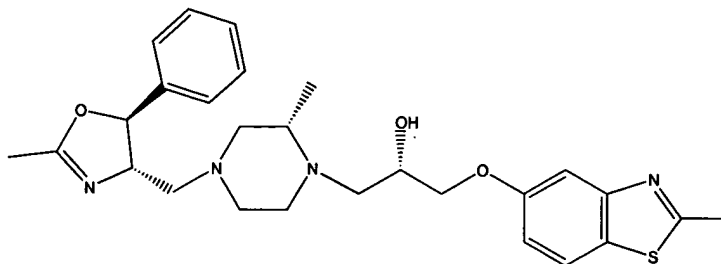
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EXAMPLE 17

Alternative Preparation of a Compound of Formula I

Preparation of a Compound of Formula I in which R¹, R², R³, R⁴, R⁵, R⁶ and R⁷ are Hydrogen, R⁸ is (S)-Methyl, T is Oxygen, X¹-Y- is 2-Methyl-5-phenyl-4,5-dihydro-oxazol-4-yl, X² is 2-Methylbenzothiazol-5-yl, and Z¹ and Z² are Methylene



- A. To a stirred mixture of ((5S)-2-methyl-5-phenyl-1,3-oxazolin-4-yl)methan-1-ol (1.0g, 5.23mmol) in methylene chloride (15mL) was added thionyl chloride (570μL, 7.81mmol) dropwise at 0°C. The resulting mixture was stirred under an atmosphere of nitrogen for 8 hours, then concentrated under a reduced pressure to afford (5S)-4-chloromethyl-2-methyl-5-phenyl-4,5-dihydro-1,3-oxazole as a white solid (960mg, M+1 = 210.1).
- B. To a stirred mixture of (2R)-3-((2S)-2-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol (51mg, 0.13mmol) in *tert*-amylalcohol (5mL) was added triethylamine (90μL, 0.65mmol), followed by (5S)-4-chloromethyl-2-methyl-5-phenyl-1,3-oxazole (29mg, 0.14mmol). The resulting mixture was stirred at 95°C for 12 hours, concentrated, and purified by preparative chromatograph (1:19 MeOH:CH₂Cl₂) to afford (2R)-3-((2S)-4-(((4S,5S)-2-methyl-5-phenyl(1,3-oxazolin-4-yl))methyl)-2-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol; (5mg, M+1 = 495.2).

B. Preparation of a Compound of Formula I

Similarly, following the procedure of 17A and 17B above, but optionally replacing ((5S)-2-methyl-5-phenyl-1,3-oxazolin-4-yl)methan-1-ol with other compounds of formula (10), and optionally replacing (2R)-3-((2S)-2-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol with other compounds of formula (6), the following compounds of Formula I were prepared:

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

5 (2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[2-chlorophenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

3-[(2S)-2-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl](2R)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

10 3-[(2S)-2-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl](2R)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-[(3S)-3-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-[(3S)-3-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

15 (2R)-3-(4-{[2-(4-fluorophenyl)(1,3-oxazolin-4-yl)]methyl}piperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

(2R)-3-((2S)-2-methyl-4-{[3-(4-methylphenyl)(4,5-dihydroisoxazol-5-yl)]methyl}piperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

20 3-(4-{[3-(4-chlorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl}(2S)-2-methylpiperazinyl)(2R)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol

(2R)-3-{4-(((4S,5S)-2-methyl-5-phenyl(1,3-oxazolin-4-yl))methyl)piperazinyl}-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;and

(2R)-3-{(2S)-4-(((4S,5S)-2-methyl-5-phenyl(1,3-oxazolin-4-yl))methyl)-2-methylpiperazinyl}-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol.

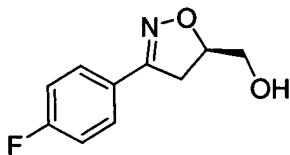
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EXAMPLE 18

Preparation of a Compound of Formula (22)

Preparation of a Compound of Formula (22) in which X¹ is 4-Fluorophenyl



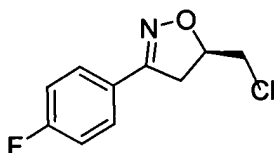
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To a cooled (0°C) solution of allyl alcohol (1.0 ml, 14.7 mmol) in chloroform (15.0 ml) was added 1.0 M diethylzinc in heptane (15.0 ml, 15.0 mmol) and stirred for 15 min under an atmosphere of nitrogen. A solution of (2*R*,3*R*)-diisopropyl 2,3-
10 dihydroxysuccinate (2.7 ml, 12.7 mmol) was added to above mixture and stirred for 2 hours. Then another portion of 1.0 M diethylzinc in heptane (15.0 ml, 15.0 mmol) was added at 0°C. A solution of α-chloro-4-fluorobenzaldoxime (2.8 g, 16.1 mmol) in chloroform (15 ml) was added to the reaction mixture. After 3 hours, the reaction was quenched with saturated NH₄Cl, and the mixture extracted with CHCl₃ (3 x 50 ml). The
15 combined organic phase was washed with brine, dried over Mg₂SO₄, filtered, and concentrated under reduced pressure to yield a crude solid. Chromatography (1:98 to 10:90 MeOH:CH₂Cl₂) over silica gel yielded ((*R*)-3-(4-fluorophenyl)-4,5-dihydroisoxazol-5-yl)methanol, a compound of formula (22) as a white solid (2.34 g, 12.0 mmol).

20

EXAMPLE 19

Preparation of (5*R*)-5-(chloromethyl)-3-(4-fluorophenyl)-4,5-dihydroisoxazole



25

Thionyl chloride (1.0 M in CH₂Cl₂, 6.0 ml, 6.0 mmol) was added dropwise to a solution of ((*R*)-3-(4-fluorophenyl)-4,5-dihydroisoxazol-5-yl)methanol (1.1 g, 5.6 mmol) in methylene chloride (10 ml) at 0°C for 1 hour. Then the reaction mixture was allowed

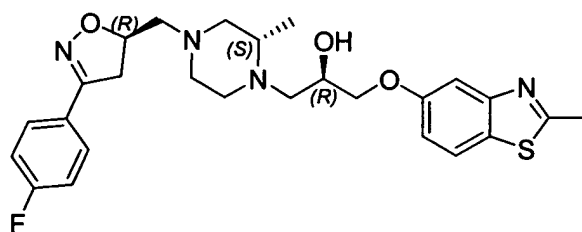
to warm to room temperature over 6 hours. The product was isolated and purified by removal of solvent under reduced pressure, followed by preparative chromatography (1:9 MeOH: CH₂Cl₂) to yield (5R)-5-(chloromethyl)-3-(4-fluorophenyl)-4,5-dihydroisoxazole (1.0 g, 4.8 mmol) as a colorless oil.

5

EXAMPLE 20

Alternative Preparation of a Compound of Formula I

10 Preparation of a Compound of Formula I in which R¹, R², R³, R⁴, R⁵, R⁶ and R⁷ are Hydrogen, R⁸ is (S)-Methyl, T is Oxygen, X¹-Y- is 3-(4-Fluorophenyl)-4,5-dihydro-oxazol-5-yl, X² is 2-Methylbenzothiazol-5-yl, and Z¹ and Z² are Methylene



15 To a mixture of (2R)-3-((2S)-2-methylpiperazinyl)-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol (6.5 mg, 0.02 mmol), and (R)-5-(chloromethyl)-3-(4-fluorophenyl)-4,5-dihydroisoxazole (4.5 mg, 0.02 mmol) in *tert*-amyl alcohol (5.0 ml) was added triethylamine (5 μ l, 0.04 mmol). The resulting reaction mixture was heated to reflux at 100°C overnight. After 16 hours the solvent was removed under reduced pressure, and
20 purified by a gradient reverse phase HPLC (acetonitrile and water as eluting solvent) to yield, after concentration (in vacuo) (2R)-1-(4-{[(5R)-3-(4-fluorophenyl)(4,5-dihydroisoxazol-5-yl)]methyl}(2S)-2-methylpiperazinyl)-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol (2.0 mg, 0.004 mmol) as a pale yellow solid.

25 The following examples illustrate the preparation of representative pharmaceutical formulations containing a compound of Formula I, such as those prepared in accordance with Examples 8 and 12.

EXAMPLE 21

Hard gelatin capsules containing the following ingredients are prepared:

	<u>Ingredient</u>	<u>Quantity</u> <u>(mg/capsule)</u>
5	Active Ingredient	30.0
	Starch	305.0
	Magnesium stearate	5.0

The above ingredients are mixed and filled into hard gelatin capsules.

10

EXAMPLE 22

A tablet formula is prepared using the ingredients below:

	<u>Ingredient</u>	<u>Quantity</u> <u>(mg/tablet)</u>
15	Active Ingredient	25.0
	Cellulose, microcrystalline	200.0
	Colloidal silicon dioxide	10.0
	Stearic acid	5.0

The components are blended and compressed to form tablets.

20

EXAMPLE 23

A dry powder inhaler formulation is prepared containing the following components:

25	<u>Ingredient</u>	<u>Weight %</u>
	Active Ingredient	5
	Lactose	95

The active ingredient is mixed with the lactose and the mixture is added to a dry powder inhaling appliance.

30

EXAMPLE 24

Tablets, each containing 30 mg of active ingredient, are prepared as follows:

	<u>Ingredient</u>	<u>Quantity</u> <u>(mg/tablet)</u>
5	Active Ingredient	30.0 mg
	Starch	45.0 mg
	Microcrystalline cellulose	35.0 mg
	Polyvinylpyrrolidone	
	(as 10% solution in sterile water)	4.0 mg
10	Sodium carboxymethyl starch	4.5 mg
	Magnesium stearate	0.5 mg
	Talc	<u>1.0 mg</u>
	Total	120 mg

The active ingredient, starch and cellulose are passed through a No. 20 mesh U.S. sieve and mixed thoroughly. The solution of polyvinylpyrrolidone is mixed with the resultant powders, which are then passed through a 16 mesh U.S. sieve. The granules so produced are dried at 50 °C to 60 °C and passed through a 16 mesh U.S. sieve. The sodium carboxymethyl starch, magnesium stearate, and talc, previously passed through a No. 30 mesh U.S. sieve, are then added to the granules which, after mixing, are compressed on a tablet machine to yield tablets each weighing 120 mg.

EXAMPLE 25

Suppositories, each containing 25 mg of active ingredient are made as follows:

25

<u>Ingredient</u>	<u>Amount</u>
Active Ingredient	25 mg
Saturated fatty acid glycerides to	2,000 mg

30

The active ingredient is passed through a No. 60 mesh U.S. sieve and suspended in the saturated fatty acid glycerides previously melted using the minimum heat

necessary. The mixture is then poured into a suppository mold of nominal 2.0 g capacity and allowed to cool.

5

EXAMPLE 26

Suspensions, each containing 50 mg of active ingredient per 5.0 mL dose are made as follows:

	<u>Ingredient</u>	<u>Amount</u>
	Active Ingredient	50.0 mg
10	Xanthan gum	4.0 mg
	Sodium carboxymethyl cellulose (11%)	
	Microcrystalline cellulose (89%)	50.0 mg
	Sucrose	1.75 g
	Sodium benzoate	10.0 mg
15	Flavor and Color	q.v.
	Purified water to	5.0 mL

The active ingredient, sucrose and xanthan gum are blended, passed through a No. 10 mesh U.S. sieve, and then mixed with a previously made solution of the microcrystalline cellulose and sodium carboxymethyl cellulose in water. The sodium benzoate, flavor, and color are diluted with some of the water and added with stirring. Sufficient water is then added to produce the required volume.

25

EXAMPLE 27

A subcutaneous formulation may be prepared as follows:

	<u>Ingredient</u>	<u>Quantity</u>
	Active Ingredient	5.0 mg
30	Corn Oil	1.0 mL

EXAMPLE 28

An injectable preparation is prepared having the following composition:

	<u>Ingredients</u>	<u>Amount</u>
5	Active ingredient	2.0 mg/ml
	Mannitol, USP	50 mg/ml
	Gluconic acid, USP	q.s. (pH 5-6)
	water (distilled, sterile)	q.s. to 1.0 ml
10	Nitrogen Gas, NF	q.s.

EXAMPLE 29

A topical preparation is prepared having the following composition:

	<u>Ingredients</u>	<u>grams</u>
15	Active ingredient	0.2-10
	Span 60	2.0
	Tween 60	2.0
	Mineral oil	5.0
20	Petrolatum	0.10
	Methyl paraben	0.15
	Propyl paraben	0.05
	BHA (butylated hydroxy anisole)	0.01
25	Water	q.s. to 100

All of the above ingredients, except water, are combined and heated to 60° C with stirring. A sufficient quantity of water at 60° C is then added with vigorous stirring to emulsify the ingredients, and water then added q.s. 100 g.

EXAMPLE 30

Sustained Release Composition

	<u>Ingredient</u>	<u>Weight Range (%)</u>	<u>Preferred Range (%)</u>	<u>Most Preferred</u>
	Active ingredient	50-95	70-90	75
	Microcrystalline cellulose (filler)	1-35	5-15	10.6
5	Methacrylic acid copolymer	1-35	5-12.5	10.0
	Sodium hydroxide	0.1-1.0	0.2-0.6	0.4
	Hydroxypropyl methylcellulose	0.5-5.0	1-3	2.0
	Magnesium stearate	0.5-5.0	1-3	2.0

The sustained release formulations of this invention are prepared as follows: compound and pH-dependent binder and any optional excipients are intimately mixed(dry-blended). The dry-blended mixture is then granulated in the presence of an aqueous solution of a strong base which is sprayed into the blended powder. The granulate is dried, screened, mixed with optional lubricants (such as talc or magnesium stearate), and compressed into tablets. Preferred aqueous solutions of strong bases are solutions of alkali metal hydroxides, such as sodium or potassium hydroxide, preferably sodium hydroxide, in water (optionally containing up to 25% of water-miscible solvents such as lower alcohols).

The resulting tablets may be coated with an optional film-forming agent, for identification, taste-masking purposes and to improve ease of swallowing. The film forming agent will typically be present in an amount ranging from between 2% and 4% of the tablet weight. Suitable film-forming agents are well known to the art and include hydroxypropyl methylcellulose, cationic methacrylate copolymers (dimethylaminoethyl methacrylate/ methyl-butyl methacrylate copolymers - Eudragit® E - Röhm. Pharma), and the like. These film-forming agents may optionally contain colorants, plasticizers, and other supplemental ingredients.

The compressed tablets preferably have a hardness sufficient to withstand 8 Kp compression. The tablet size will depend primarily upon the amount of compound in the tablet. The tablets will include from 300 to 1100 mg of compound free base. Preferably,

the tablets will include amounts of compound free base ranging from 400-600 mg, 650-850 mg, and 900-1100 mg.

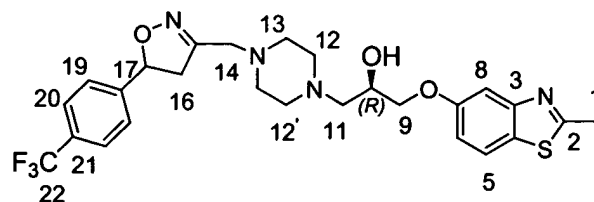
In order to influence the dissolution rate, the time during which the compound containing powder is wet mixed is controlled. Preferably the total powder mix time, i.e. the time during which the powder is exposed to sodium hydroxide solution, will range from 1 to 10 minutes and preferably from 2 to 5 minutes. Following granulation, the particles are removed from the granulator and placed in a fluid bed dryer for drying at about 60°C.

EXAMPLE 31 Mitochondrial Assays

Rat heart mitochondria were isolated by the method of Nedergard and Cannon (Methods in Enzymol. 55, 3, 1979).

Palmitoyl CoA oxidation – The Palmitoyl CoA oxidation was carried out in a total volume of 100 micro liters containing the following agents: 110 mM KCl, 33 mM Tris buffer at pH 8, 2 mM KPi, 2 mM MgCl₂, 0.1 mM EDTA, 14.7 microM defatted BSA, 0.5 mM malic acid, 13 mM carnitine, 1 mM ADP, 52 micrograms of mitochondrial protein, and 16 microM 1-C¹⁴ palmitoyl CoA (Sp. Activity 60 mCi/mmol; 20 microCi/ml, using 5 microliters per assay). The compounds of this invention were added in a DMSO solution at the following concentrations: 100 micro molar, 30 micro molar, and 3 micro molar. In each assay, a DMSO control was used. After 15 min at 30°C, the enzymatic reaction was centrifuged (20,000 g for 1 min), and 70 microliters of the supernatant was added to an activated reverse phase silicic acid column (approximately 0.5 ml of silicic acid). The column was eluted with 2 ml of water, and 0.5 ml of the eluent was used for scintillation counting to determine the amount of C¹⁴ trapped as C¹⁴ bicarbonate ion.

The compounds of the invention showed activity as fatty acid oxidation inhibitors in this assay. Representative examples of test data is shown below, along with the NMR determined for the compound.

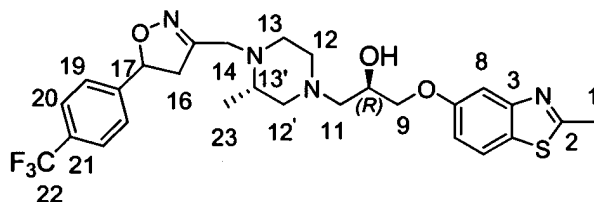


(2R)-1-(2-methylbenzothiazol-5-yloxy)-3-[4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]propan-2-ol;

1HNMR (400 MHz CD₃OD) δ 7.95 (d, J = 9.00 Hz, 1H, H5), 7.73 (d, J = 8.22 Hz, 2H, H20), 7.61 (d, J = 8.22 Hz, 2H, H19), 7.52 (d, J = 2.35 Hz, 1H, H8), 7.26 (dd, J = 2.74, 9.00 Hz, 1H, H6), 5.82 (dd, J = 8.22, 11.4 Hz, 1H, H17), 4.53 (m, 1H, H10), 4.18 (d, J = 4.70 Hz, 2H, H9), 3.91 (s, 2H, H14), 3.47-3.76 (m, 6H, H11, H12, H12', H13, H13'), 3.12 (m, 2H, H16), 2.97 (s, 3H, H1); MS: m/z 535 (100, MH⁺).

EC₅₀ = 1.70 mM, Liver S-9 index: 1.00b

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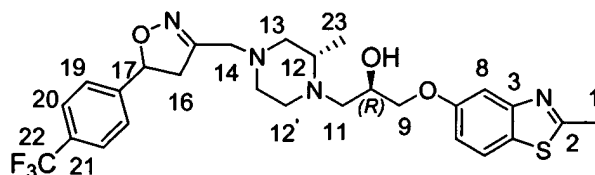


(2R)-3-[(3S)-3-methyl-4-({5-[4-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-1-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

1HNMR (400 MHz CD₃OD) δ 7.74 (m, 1H, H5), 7.65 (m, 2H, H20), 7.52 (m, 2H, H19), 7.40 (m, 1H, H8), 7.06 (m, 1H, H6), 5.67 (m, 1H, H17), 4.13 (m, 1H, H10), 4.05 (m, 2H, H9), 2.82-3.63 and 2.30-2.70 (m, 11H, H14, H11, H12, H12', H13, H13'), 2.78 (s, 3H, H1), 2.05 (m, 2H, H16), 1.08 (d, 3H, H23); MS: m/z 549 (100, MH⁺).

EC₅₀ = 0.24 mM, Liver S-9 index: 0.39b

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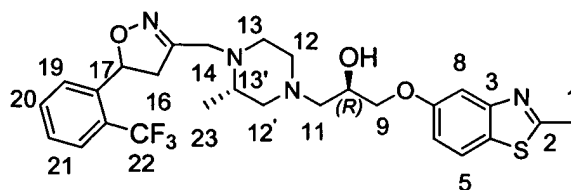


(2R)-1-[(2S)-2-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

1HNMR (400 MHz CD₃OD) δ 7.70 (d, J = 9.00 Hz, 1H, H₅), 7.75 (d, J = 8.23 Hz, 2H, H₂₀), 7.64 (d, J = 8.23 Hz, 2H, H₁₉), 7.44 (d, J = 2.33 Hz, 1H, H₈), 7.20 (dd, J = 2.73, 9.00 Hz, 1H, H₆), 5.69 (dd, J = 8.18, 10.9 Hz, 1H, H₁₇), 4.44 (m, 1H, H₁₀), 4.14 (d, J = 4.72 Hz, 2H, H₉), 3.83 (s, 2H, H₁₄), 3.40-3.71 (m, 6H, H₁₁, H₁₂, H_{12'}, H₁₃, H_{13'}), 3.08 (m, 2H, H₁₆), 2.91 (s, 3H, H₁); 1.28 (d, 3H, H₂₃); MS: m/z 549 (100, MH⁺).

EC₅₀ = 0.14 mM, Liver S-9 index: NA

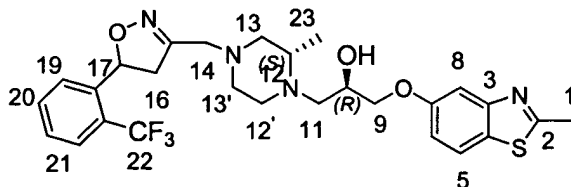
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(2R)-1-[(3S)-3-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

1HNMR (400 MHz CD₃OD) δ 7.87 (m, 1H, H₅), 7.70 (m, 2H, H₂₀), 7.55 (m, 2H, H₁₉), 7.41 (m, 1H, H₈), 7.08 (m, 1H, H₆), 5.60 (m, 1H, H₁₇), 4.16 (m, 1H, H₁₀), 4.08 (m, 2H, H₉), 2.78-3.60 and 2.28-2.67 (m, 11H, H₁₄, H₁₁, H₁₂, H_{12'}, H₁₃, H_{13'}), 2.72 (s, 3H, H₁), 2.00 (m, 2H, H₁₆), 1.13 (d, 3H, H₂₃); MS: m/z 549 (100, MH⁺).

EC₅₀ = 0.17 mM, Liver S-9 index: 0.03b.

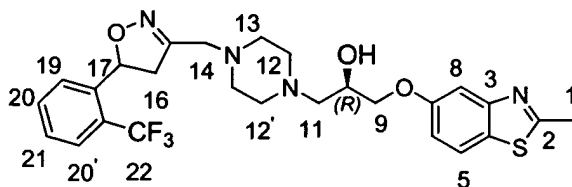


(2R)-1-[(2S)-2-methyl-4-({5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl)}methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

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¹HNMR (400 MHz CD₃OD) δ 7.72 (m, 1H, H₅), 7.58 (m, 3H, H₁₉, H₂₀, H_{20'}), 7.47 (t, J = 7.83 Hz, 1H, H₂₁), 7.38 (m, 1H, H₈), 7.10 (m, 1H, H₆), 5.63 (m, 1H, H₁₇), 4.49 (m, 1H, H₁₀), 4.17 (m, 2H, H₉), 3.84 (s, 2H, H₁₄), 3.42-3.75 (m, 6H, H₁₁, H₁₂, H_{12'}, H₁₃, H_{13'}), 3.09 (m, 2H, H₁₆), 2.87 (s, 3H, H₁); 1.20 (d, 3H, H₂₃); MS: m/z 549 (100, MH⁺).

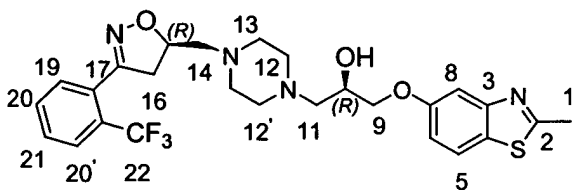
5 EC₅₀ = 0.40 mM, Liver S-9 index: NA



(2R)-3-(2-methylbenzothiazol-5-yloxy)-1-[4-((5-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-3-yl))methyl)piperazinyl]propan-2-ol;

10 ¹HNMR (400 MHz CD₃OD) δ 7.91 (d, J = 9.00 Hz, 1H, H₅), 7.68 (m, 3H, H₁₉, H₂₀, H_{20'}), 7.51 (t, J = 7.80 Hz, 1H, H₂₁), 7.47 (d, J = 2.34 Hz, 1H, H₈), 7.21 (dd, J = 2.74, 9.00 Hz, 1H, H₆), 6.01 (dd, J = 8.61, 10.28 Hz, 1H, H₁₇), 4.50 (m, 1H, H₁₀), 4.14 (d, J = 4.70 Hz, 2H, H₉), 3.84 (s, 2H, H₁₄), 3.42-3.74 (m, 6H, H₁₁, H₁₂, H_{12'}, H₁₃, H_{13'}), 3.00 (m, 2H, H₁₆), 2.92 (s, 3H, H₁); MS: m/z 535 (100, MH⁺).

15 EC₅₀ = 1.74 mM, Liver S-9 index: NA



(2R)-1-[4-((5R)-3-[2-(trifluoromethyl)phenyl](4,5-dihydroisoxazol-5-yl))methyl)piperazinyl]-3-(2-methylbenzothiazol-5-yloxy)propan-2-ol;

20 ¹HNMR (400 MHz CD₃OD) δ 7.98 (d, J = 9.00 Hz, 1H, H₅), 7.70 (m, 3H, H₁₉, H₂₀, H_{20'}), 7.53 (t, J = 7.93 Hz, 1H, H₂₁), 7.50 (d, J = 2.36 Hz, 1H, H₈), 7.23 (dd, J = 2.74, 9.00 Hz, 1H, H₆), 4.52 (m, 1H, H₁₀), 4.18 (d, J = 4.69 Hz, 2H, H₉), 3.90 (m, 1H, H₁₅), 3.83 (s, 2H, H₁₄), 3.35-3.70 (m, 6H, H₁₁, H₁₂, H_{12'}, H₁₃, H_{13'}), 2.97 (m, 2H, H₁₆), 2.90 (s, 3H, H₁); MS: m/z 535 (100, MH⁺).

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IC50 values are the mean of n=3, experiments were carried out at 6 different concentrations. Each experiment was carried out in triplicate. Experiments were carried out using human liver S-9.

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EXAMPLE 32

Perfusate

Langendorff perfusion is conducted using a Krebs-Henseleit solution containing: (mM) NaCl (118.0), KCl (4.7), KH₂PO₄ (1.2), MgSO₄ (1.2), CaCl₂ (2.5), NaHCO₃ (25.0) and
10 glucose (5.5 or 11) (Finegan et al. 1996). The working heart perfusate consists of a Krebs-Henseleit solution with the addition of palmitate (0.4 or 1.2 mM) pre-bound to 3% bovine serum albumin (essentially fatty acid free BSA) and insulin (100 µU/ml). Palmitate is initially dissolved in an ethanol:water mixture (40%:60%) containing 0.5-0.6 g Na₂CO₃ per g of palmitate. Following heating to evaporate the ethanol, this mixture is
15 then added to the 3% BSA-Krebs-Henseleit mixture (without glucose) and allowed to dialyze (8000 MW cut-off) overnight in 10 volumes of glucose-free Krebs-Henseleit solution. The next day, glucose is added to the solution and the mixture was filtered through glass microfiber filters (GF/C, Whatman, Maidstone, England) and kept on ice, or refrigerated, prior to use. The perfusate is continuously oxygenated with a 95% CO₂,
20 5% O₂ gas mixture while in the perfusion apparatus to main aerobic conditions.

EXAMPLE 33

Heart Perfusion Protocols

25 Rats are anesthetized with pentobarbital (60 mg/kg, intraperitoneally) and hearts rapidly removed and placed in ice-cold Krebs-Henseleit solution. The hearts are then rapidly cannulated via the aortic stump and Langendorff perfusion at constant pressure (60 mm Hg) is initiated and continued for a 10-min equilibration period. During this equilibration period, the pulmonary artery is cut, and excess fat and lung tissue removed
30 to reveal the pulmonary vein. The left atrium is cannulated and connected to the preload line originating from the oxygenation chamber. After the 10-min equilibration period,

hearts are switched to working mode (by clamping off the Langendorff line and opening the preload and afterload lines) and perfused at 37°C under aerobic conditions at a constant left atrial preload (11.5 mm Hg) and aortic afterload (80 mm Hg). The compliance chamber is filled with air adequate to maintain developed pressure at 50-60 mm Hg. Perfusate is delivered to the oxygenation chamber via a peristaltic pump from the reservoir chamber that collected aortic and coronary flows as well as overflow from the oxygenator.

Typically, hearts are perfused under aerobic conditions for 60 min. Hearts are paced at 300 beats/min throughout each phase of the perfusion protocol (voltage adjusted as necessary) with the exception of the initial 5 min of reperfusion when hearts are allowed to beat spontaneously.

At the end of the perfusion protocol, hearts are rapidly frozen using Wollenberger clamps cooled to the temperature of liquid nitrogen. Frozen tissues are pulverized and the resulting powders stored at -80°C.

EXAMPLE 34

Myocardial Mechanical Function

Aortic systolic and diastolic pressures are measured using a Sensoror (Horten Norway) pressure transducer attached to the aortic outflow line and connected to an AD Instruments data acquisition system. Cardiac output, aortic flow and coronary flow (cardiac output minus aortic flow) are measured (ml/min) using in-line ultrasonic flow probes connected to a Transonic T206 ultrasonic flow meter. Left ventricular minute work (LV work), calculated as cardiac output x left ventricular developed pressure (aortic systolic pressure - preload pressure), is used as a continuous index of mechanical function. Hearts are excluded if LV work decreased more than 20% during the 60-min period of aerobic perfusion.

EXAMPLE 35

Myocardial Oxygen Consumption and Cardiac Efficiency

Measuring the atrial-venous difference in oxygen content of the perfusate and multiplying by the cardiac output provides an index of oxygen consumption. Atrial oxygen content (mmHg) is measured in perfusate in the preload line or just prior to entering the left atria. Venous oxygen content is measured from perfusate exiting the pulmonary artery and passing through in-line O₂ probes and meters Microelectrodes Inc., Bedford, NH. Cardiac efficiency is calculated as the cardiac work per oxygen consumption.

EXAMPLE 36

Measurement of Glucose and Fatty Acid Metabolism

Determining the rate of production of ³H₂O and ¹⁴CO₂ from [³H/¹⁴C]glucose in the isolated working rat model allows a direct and continuous measure of the rates of glycolysis and glucose oxidation. Alternatively, the measure of the production of ³H₂O from [5-³H]palmitate provides a direct and continuous measure of the rate of palmitate oxidation. Dual labelled substrates allows for the simultaneous measure of either glycolysis and glucose oxidation or fatty acid oxidation and glucose oxidation. A 3-ml sample of perfusate is taken from the injection port of the recirculating perfusion apparatus at various time-points throughout the protocol for analysis of ³H₂O and ¹⁴CO₂ and immediately placed under mineral oil until assayed for metabolic product accumulation. Perfusate is supplemented with [³H/¹⁴C]glucose or [5-³H]palmitate to approximate a specific activity of 20 dpm/mmol. Average rates of glycolysis and glucose oxidation are calculated from linear cumulative time-courses of product accumulation between 15 and 60 min for aerobic perfusion. Rates of glycolysis and glucose oxidation are expressed as μmol glucose metabolized/min/g dry wt.

EXAMPLE 37

Measurement of Myocardial Glycolysis

Rates of glycolysis are measured directly as previously described (Saddik & Lopaschuk, 1991) from the quantitative determination of $^3\text{H}_2\text{O}$ liberated from radiolabeled $[5\text{-}^3\text{H}]\text{glucose}$ at the enolase step of glycolysis. Perfusate samples are collected at various time-points throughout the perfusion protocol. $^3\text{H}_2\text{O}$ is separated from the perfusate by passing perfusate samples through columns containing Dowex 1-X 4 anion exchange resin (200-400 mesh). A 90 g/L Dowex in 0.4 M potassium tetraborate mixture is stirred overnight after which 2 ml of the suspension is loaded into separation columns and washed extensively with dH_2O to remove the tetraborate. The columns are found to exclude 98-99.6 % of the total $[^3\text{H}]\text{glucose}$ (Saddik & Lopaschuk, 1996). Perfusate samples (100 μl) are loaded onto the columns and washed with 1.0 ml dH_2O . Effluent is collected into 5 ml of Ecolite Scintillation Fluid (ICN, Radiochemicals, Irvine, CA) and counted for 5 min in a Beckman LS 6500 Scintillation Counter with an automatic dual ($^3\text{H}/^{14}\text{C}$) quench correction program. Average rates of glycolysis for each phase of perfusion are expressed as μmol glucose metabolized/min/g dry wt as described above.

EXAMPLE 38

Measurement of Myocardial Glucose Oxidation

Glucose oxidation is also determined directly as previously described (Saddik & Lopaschuk, 1991) by measuring $^{14}\text{CO}_2$ from $[^{14}\text{C}]\text{glucose}$ liberated at the level of pyruvate dehydrogenase and in the Krebs cycle. Both $^{14}\text{CO}_2$ gas exiting the oxygenation chamber and $[^{14}\text{C}]\text{bicarbonate}$ retained in solution are measured. Perfusate samples are collected at various time-points throughout the perfusion protocol. $^{14}\text{CO}_2$ gas is collected by passing the gas exiting the oxygenator through a hyamine hydroxide trap (20-50 ml depending on perfusion duration). Perfusate samples (2 x 1 ml), which are stored under oil to prevent the escape of gas by equilibration with atmospheric CO_2 , are injected into 16 x 150 mm test tubes containing 1 ml of 9 N H_2SO_4 . This process releases $^{14}\text{CO}_2$ from the perfusate present as $\text{H}^{14}\text{CO}_3^-$. These duplicate tubes are sealed with a rubber stopper

attached to a 7-ml scintillation vial containing a 2 x 5 cm piece of filter paper saturated with 250 μ l of hyamine hydroxide. The scintillation vials with filter papers are then removed and Ecolite Scintillation Fluid (7 ml) added. Samples are counted by standard procedures as described above. Average rates of glucose oxidation for each phase of perfusion are expressed as μ mol glucose metabolized/min/g dry wt as described above.

EXAMPLE 39

Measurement of Myocardial Fatty Acid Oxidation

Rates of palmitate oxidation are measured directly as previously described (Saddik & Lopaschuk, 1991) from the quantitative determination of $^3\text{H}_2\text{O}$ liberated from radiolabeled $[5-^3\text{H}]\text{palmitate}$. $^3\text{H}_2\text{O}$ is separated from $[5-^3\text{H}]\text{palmitate}$ following a chloroform:methanol (1.88 ml of 1:2 v/v) extraction of a 0.5 ml sample of buffer then adding 0.625 ml of chloroform and 0.625 ml of a 2M KCL:HCl solution. The aqueous phase is removed and treated with a mixture of chloroform, methanol and KCl:HCl (1:1:0.9 v/v). Duplicate samples are taken from the aqueous phase for liquid scintillation counting and rates of oxidation are determined taking into account a dilution factor. This results in >99% extraction and separation of $^3\text{H}_2\text{O}$ from $[5-^3\text{H}]\text{palmitate}$. Average rates of glucose oxidation for each phase of perfusion are expressed as μ mol glucose metabolized/min/g dry wt as described above.

Dry to Wet Ratios

Frozen ventricles are pulverized at the temperature of liquid nitrogen with a mortar and pestle. Dry to wet determinations are made by weighing a small amount of frozen heart tissue and re-weighing that same tissue after 24-48 hr of air drying and taking the ratio of the two weights. From this ratio, total dry tissue can be calculated. This ratio is used to normalize, on a per g dry weight basis, rates of glycolysis, glucose oxidation and glycogen turnover as well as metabolite contents.

The compounds of the invention show activity as fatty acid oxidation inhibitors in this assay.

REFERENCES

1. Finegan BA, Gandhi M, Lopaschuk GD, Clanachan AS, 1996. Antecedent ischemia reverses effects of adenosine on glycolysis and mechanical function of working hearts. *American Journal of Physiology* 271: H2116-25.

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2. Saddik M, Lopaschuk GD, 1991. Myocardial triglyceride turnover and contribution to energy substrate utilization in isolated working rat hearts. *Journal of Biological Chemistry* 266: 8162-8170.

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While the present invention has been described with reference to the specific embodiments thereof, it will be understood by those skilled in the art that various changes may be made and equivalents may be substituted without departing from the true spirit and scope of the invention. In addition, many modifications may be made to adapt a particular situation, material, composition of matter, process, process step or steps, to the objective, spirit and scope of the present invention. All such modifications are intended to be within the scope of the claims appended hereto. All patents and publications cited above are hereby incorporated by reference.

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